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INHIBITORS OF ASPARTYL PROTEASE

TECHNICAL FIELD OF THE INVENTION

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The present invention relates to a novel class of sulfonamides which are aspartyl protease inhibitors. In one embodiment, this invention relates to a novel class of HIV aspartyl protease inhibitors characterized by specific structural and physicochemical features. This invention also relates to pharmaceutical compositions comprising these compounds. The compounds and pharmaceutical compositions of this invention are particularly well suited for inhibiting HIV-1 and HIV-2 protease activity and consequently, may be advantageously used as anti-viral agents against the HIV-1 and HIV-2 viruses. This invention also relates to methods for inhibiting the activity of HIV aspartyl protease using the compounds of this invention and methods for screening compounds for anti-HIV activity.

BACKGROUND OF THE INVENTION

The human immunodeficiency virus ("HIV") is the causative agent for acquired immunodeficiency syndrome ("AIDS") -- a disease characterized by the destruction of the immune system, particularly of CD4⁺ T-cells, with

attendant susceptibility to opportunistic infections -- and its precursor AIDS-related complex ("ARC") -- a syndrome characterized by symptoms such as persistent generalized lymphadenopathy, fever and weight loss.

5 As in the case of several other retroviruses, HIV encodes the production of a protease which carries out post-translational cleavage of precursor polypeptides in a process necessary for the formation of infectious virions (S. Crawford et al., "A Deletion Mutation in the
10 5' Part of the pol Gene of Moloney Murine Leukemia Virus Blocks Proteolytic Processing of the gag and pol Polyproteins", J. Virol., 53, p. 899 (1985)). These gene products include pol, which encodes the virion RNA-dependent DNA polymerase (reverse transcriptase), an
15 endonuclease, HIV protease, and gag, which encodes the core-proteins of the virion (H. Toh et al., "Close Structural Resemblance Between Putative Polymerase of a Drosophila Transposable Genetic Element 17.6 and pol gene product of Moloney Murine Leukemia Virus", EMBO J., 4, p. 1267 (1985); L.H. Pearl et al., "A Structural Model
20 for the Retroviral Proteases", Nature, pp. 329-351 (1987); M.D. Power et al., "Nucleotide Sequence of SRV-1, a Type D Simian Acquired Immune Deficiency Syndrome Retrovirus", Science, 231, p. 1567 (1986)).

25 A number of synthetic anti-viral agents have been designed to target various stages in the replication cycle of HIV. These agents include compounds which block viral binding to CD4⁺ T-lymphocytes (for example, soluble CD4), and compounds which interfere with viral
30 replication by inhibiting viral reverse transcriptase (for example, didanosine and zidovudine (AZT)) and inhibit integration of viral DNA into cellular DNA (M.S. Hirsh and R.T. D'Aquila, "Therapy for Human

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Immunodeficiency Virus Infection", N.Eng.J.Med., 328, p. 1686 (1993)). However, such agents, which are directed primarily to early stages of viral replication, do not prevent the production of infectious virions in chronically infected cells. Furthermore, administration of some of these agents in effective amounts has led to cell-toxicity and unwanted side effects, such as anemia and bone marrow suppression.

More recently, the focus of anti-viral drug design has been to create compounds which inhibit the formation of infectious virions by interfering with the processing of viral polyprotein precursors. Processing of these precursor proteins requires the action of virus-encoded proteases which are essential for replication (Kohl, N.E. et al. "Active HIV Protease is Required for Viral Infectivity" Proc. Natl. Acad. Sci. USA, 85, p. 4686 (1988)). The anti-viral potential of HIV protease inhibition has been demonstrated using peptidyl inhibitors.

More recently several small molecule protease inhibitors have become available for the treatment of HIV infections. Among these is the sulfonamide-containing molecule, Agenerase® (amprenavir). Agenerase® is described in United States patent 5,585,397. Other sulfonamide inhibitors of aspartyl protease are described in United States patents 5,691,372, 5,510,388, 5,521,219, 5,639,769, 5,714,605, 5,744,481, 5,786,483, 5,830,897 and 5,843,946.

Because HIV-infected patients often develop resistance to particular protease inhibitors, the need still exists for additional compounds that can effectively inhibit the action of aspartyl proteases, particularly HIV protease, for use as agents for

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preventing and treating chronic and acute viral infections. Moreover, there is also a need for compounds that can effectively inhibit the action of HIV mutants which are resistant to conventional protease inhibitors.

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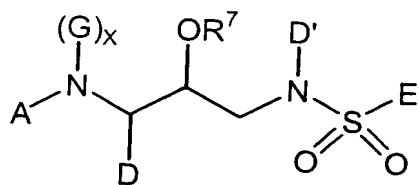
SUMMARY OF THE INVENTION

The present invention provides a novel class of compounds, and pharmaceutically acceptable derivatives thereof, that are useful as inhibitors of aspartyl proteases, in particular, HIV aspartyl protease. These compounds can be used alone or in combination with other therapeutic or prophylactic agents, such as anti-virals, antibiotics, immunomodulators or vaccines, for the treatment or prophylaxis of viral infection.

According to a preferred embodiment, the compounds of this invention are capable of inhibiting HIV viral replication in human CD₄⁺ T-cells. These compounds are useful as therapeutic and prophylactic agents to treat or prevent infection by HIV-1 and related viruses which may result in asymptomatic infection, AIDS-related complex ("ARC"), acquired immunodeficiency syndrome ("AIDS"), or similar disease of the immune system.

According to another preferred embodiment, the compounds of this invention are useful as therapeutic and prophylactic agents to treat or prevent infection by HIV mutants.

It is a principal object of this invention to provide a novel class of sulfonamides which are aspartyl protease inhibitors, and particularly, HIV aspartyl protease inhibitors. The novel sulfonamides of this invention are those of formula I:



(I)

and pharmaceutically acceptable salts thereof;
wherein:

A is selected from H; Ht; $-\text{R}^1\text{-Ht}$; $-\text{R}^1\text{-C}_1\text{-C}_6$ alkyl,
5 which is optionally substituted with one or more groups
independently selected from hydroxy, $-\text{CN}$, $\text{C}_1\text{-C}_4$ alkoxy,
Ht, $-\text{O-Ht}$, $-\text{NR}^2\text{-Ht}$, $-\text{NR}^2\text{-CO-N(R}^2)_2$, $-\text{SO}_2\text{-N(R}^2)_2$, $-\text{SO}_2\text{-R}^2$ or
 $-\text{CO-N(R}^2)_2$; $-\text{R}^1\text{-C}_2\text{-C}_6$ alkenyl, which is optionally
substituted with one or more groups independently
10 selected from hydroxy, $\text{C}_1\text{-C}_4$ alkoxy, Ht, $-\text{O-Ht}$,
 $-\text{NR}^2\text{-CO-N(R}^2)_2$ or $-\text{CO-N(R}^2)_2$; or R^7 ;

each R^1 is independently selected from $-\text{C(O)-}$,
 $-\text{S(O)}_2\text{-}$, $-\text{C(O)-C(O)-}$, $-\text{O-C(O)-}$, $-\text{O-S(O)}_2\text{-}$, $-\text{NR}^2\text{-}$,
 $-\text{NR}^2\text{-S(O)}_2\text{-}$, $-\text{NR}^2\text{-C(O)-}$ or $-\text{NR}^2\text{-C(O)-C(O)-}$;

15 each Ht is independently selected from $\text{C}_3\text{-C}_7$
cycloalkyl; $\text{C}_5\text{-C}_7$ cycloalkenyl; $\text{C}_6\text{-C}_{14}$ aryl; or a 5-7
membered saturated or unsaturated heterocycle, containing
one or more heteroatoms selected from N, $\text{N(R}^2)$, O, S and
 S(O)_n ; wherein said aryl or said heterocycle is optionally
20 fused to Q; and wherein any member of said Ht is
optionally substituted with one or more substituents
independently selected from oxo, $-\text{OR}^2$, SR^2 , $-\text{R}^2$,
 $-\text{N(R}^2)(\text{R}^2)$, $-\text{R}^2\text{-OH}$, $-\text{CN}$, $-\text{CO}_2\text{R}^2$, $-\text{C(O)-N(R}^2)_2$, $-\text{S(O)}_2\text{-N(R}^2)_2$,
 $-\text{N(R}^2)\text{-C(O)-R}^2$, $-\text{N(R}^2)\text{-C(O)O-R}^2$, $-\text{C(O)-R}^2$, $-\text{S(O)}_n\text{-R}^2$, $-\text{OCF}_3$,
25 $-\text{S(O)}_n\text{-Q}$, methylenedioxy, $-\text{N(R}^2)\text{-S(O)}_2(\text{R}^2)$, halo, $-\text{CF}_3$,
 $-\text{NO}_2$, Q, $-\text{OQ}$, $-\text{OR}^7$, $-\text{SR}^7$, $-\text{R}^7$, $-\text{N(R}^2)(\text{R}^7)$ or $-\text{N(R}^7)_2$;

each R^2 is independently selected from H, or $\text{C}_1\text{-C}_4$
alkyl optionally substituted with a 3-7 membered
saturated, partially saturated or unsaturated carbocyclic
30 ring system; or a 5-7 membered saturated, partially

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saturated or unsaturated heterocyclic ring containing one or more heteroatoms selected from O, N, S, S(O)_n or N(R³³); wherein any of said ring systems or N(R³³) is optionally substituted with 1 to 4 substituents

5 independently selected from -X'-Y', -O-arylalkyl, -S-arylalkyl, -N(Y')₂, -N(H)-arylalkyl, -N(C₁-C₄ alkyl)-arylalkyl, oxo, -O-(C₁-C₄ alkyl), OH, C₁-C₄ alkyl, -SO₂H, -SO₂-(C₁-C₄ alkyl), -SO₂-NH₂, -SO₂-NH(C₁-C₄ alkyl), -SO₂-N(C₁-C₄ alkyl)₂, -NH₂, -NH(C₁-C₄ alkyl), -N(C₁-C₄ alkyl)₂, -NH-C(O)H, -N(C₁-C₄ alkyl)-C(O)H, -NH-C(O)-C₁-C₄ alkyl, -C₁-C₄ alkyl-OH, -OH, -CN, -C(O)OH, -C(O)O-C₁-C₄ alkyl, -C(O)-NH₂, -C(O)-NH(C₁-C₄ alkyl), -C(O)-N(C₁-C₄ alkyl)₂, halo or -CF₃;

X' is -O-, -S-, -NH-, -NHC(O)-, -NHC(O)O-, -NHSO₂-,

15 or -N(C₁-C₄)alkyl-;

Y' is C₁-C₁₅ alkyl, C₂-C₁₅ alkenyl or alkynyl, wherein one to five carbon atoms in Y are optionally substituted with C₃-C₇ cycloalkyl or C₅-C₆ cycloalkenyl, C₆-C₁₄ aryl or a 5-7 membered saturated or unsaturated heterocycle,

20 containing one or more heteroatoms selected from N, NH, O, S and S(O)_n;

each R³ is independently selected from H, Ht, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₆ cycloalkyl or C₅-C₆ cycloalkenyl; wherein any member of said R³, except

25 H, is optionally substituted with one or more substituents selected from -OR², -C(O)-N(R²)₂, -S(O)_n-N(R²)₂, -N(R²)₂, -N(R²)-C(O)O(R²), -N(R²)-C(O)N(R²)₂, -N(R²)-C(O)-R², Ht, -CN, -SR², -C(O)OR², N(R²)-C(O)-R²;

each R³³ is selected from H, C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₆ cycloalkyl or C₅-C₆ cycloalkenyl, C₆-C₁₄ aryl or a 5-7 membered saturated or unsaturated heterocycle, containing one or more heteroatoms selected from N, NH, O, S and S(O)_n;

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each n is independently 1 or 2;

G, when present, is selected from H, R⁷ or C₁-C₄ alkyl, or, when G is C₁-C₄ alkyl, G and R⁷ are bound to one another either directly or through a C₁-C₃ linker to form
5 a heterocyclic ring; or

when G is not present (i.e., when x in (G)_x is 0), then the nitrogen to which G is attached is bound directly to the R⁷ group in -OR⁷ with the concomitant displacement of one -ZM group from R⁷;

10 D is selected from C₁-C₆ alkyl which is substituted with Q, which is optionally substituted with one or more groups selected from C₃-C₆ cycloalkyl, -R³, -O-Q or Q; C₂-C₄ alkenyl which is substituted with Q, which is optionally substituted with one or more groups selected
15 from -OR², -S-Ht, -R³, -O-Q or Q; C₃-C₆ cycloalkyl, which is optionally substituted with or fused to Q; or C₅-C₆ cycloalkenyl, which is optionally substituted with or fused to Q;

each Q is independently selected from a 3-7 membered
20 saturated, partially saturated or unsaturated carbocyclic ring system; or a 5-7 membered saturated, partially saturated or unsaturated heterocyclic ring containing one or more heteroatoms selected from O, N, S, S(O)_n or N(R²); wherein Q contains one substituent selected from -OR², -
25 OR⁸, -O-arylalkyl, -SR⁸, -S-arylalkyl, -N(R²)R⁸, -N(R²)-arylalkyl and may be optionally substituted with one or more additional substituents independently selected from oxo, -OR⁸, -O-arylalkyl -SR⁸, -S-arylalkyl, -N(R²)R⁸, -N(R²)-arylalkyl, -OR², -R², -SO₂R², -SO₂-N(R²)₂, -N(R²)₂,
30 -N(R²)-C(O)-R², -OH, (C₁-C₄)-OH, -CN, -CO₂R², -C(O)-N(R²)₂, halo or -CF₃;

each R⁸ is independently selected from Ht, -C₁-C₁₅ branched or straight chain alkyl, alkenyl or alkynyl

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wherein one to five carbon atoms in said alkyl, alkenyl or alkynyl are independently replaced by W, or wherein one to five carbon atoms in said alkyl, alkenyl or alkynyl are substituted with Ht; and wherein R⁸ is
5 additionally and optionally substituted with one or more groups independently selected from -OH, -S(C₁-C₆ alkyl), -CN, -CF₃, -N(R²)₂, halo, -C₁-C₄-alkyl, -C₁-C₄-alkoxy; -Ht; -O-Ht; -NR²-CO-N(R²)₂; -CO-N(R²)₂; -R¹-C₂-C₆ alkenyl, which
10 is optionally substituted with one or more groups independently selected from hydroxy, C₁-C₄ alkoxy, Ht, -O-Ht, -NR²-CO-N(R²)₂ or -CO-N(R²)₂; or R⁷;

wherein W is -O-, -NR²-, -S-, -C(O)-, -C(S)-, -C(=NR²)-, -S(O)₂-, -NR²-S(O)₂-, -S(O)₂-NR²-, -NR²-C(O)O-, -O-C(O)NR²-, -NR²-C(O)NR²-, -NR²-C(S)NR²-, -CONR², -NR²C(O)-, -C(S)NR², -NR²C(S)-, -NR²-C(=N-CN)-NR²-, -NR²C(=N-CN)O- or -C(O)O-;

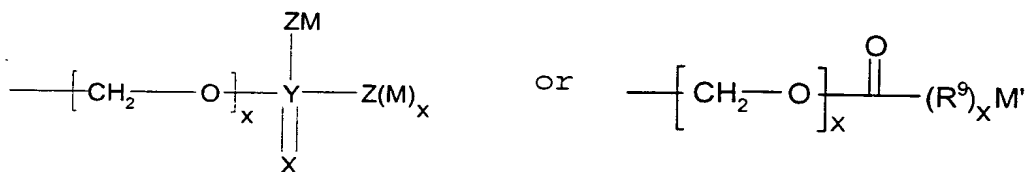
D' is selected from C₁-C₁₅ alkyl, C₁-C₁₅ alkoxy, C₂-C₁₅ alkenyl, C₂-C₁₅ alkenyloxy, C₂-C₁₅ alkynyl, or C₂-C₁₅ alkynyloxy, wherein D' optionally comprises one or more
20 substituents independently selected from Ht, oxo, halo, -CF₃, -OCF₃, -NO₂, azido, -SH, -SR³, -N(R³)-N(R³)₂, -O-N(R³)₂, -(R³)N-O-(R³), -N(R³)₂, -CN, -CO₂R³, -C(O)-N(R³)₂, -S(O)_n-N(R³)₂, -N(R³)-C(O)-R³, -N(R³)-C(O)-N(R³)₂, -C(O)-R³, -S(O)_n-R³, -N(R³)-S(O)_n(R³), -N(R³)-S(O)_n-N(R³)₂,
25 -S-NR³-C(O)R³, -C(S)N(R³)₂, -C(S)R³, -NR³-C(O)OR³, -O-C(O)OR³, -O-C(O)N(R³)₂, -NR³-C(S)R³, =N-OH, =N-OR³, =N-N(R³)₂, =NR³, =NNR³C(O)N(R³)₂, =NNR³C(O)OR³, =NNR³S(O)_n-N(R³)₂, -NR³-C(S)OR³, -NR³-C(S)N(R³)₂, -NR³-C[=N(R³)]-N(R³)₂, -N(R³)-C[=N-NO₂]-N(R³)₂,
30 -N(R³)-C[=N-NO₂]-OR³, -OC(O)R³, -OC(S)R³, -OC(O)N(R³)₂, -C(O)N(R³)-N(R³)₂, -N(R³)-N(R³)C(O)R³, -N(R³)-OC(O)R³, -N(R³)-OC(O)R³, -N(R³)-OC(O)R³, -OC(S)N(R³)₂, -OC(S)N(R³)(R³), or -PO₃-R³;

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E is selected from Ht; O-Ht; Ht-Ht; Ht fused with Ht; -O-R³; -N(R²)(R³); -N(R²)-Ht; C₁-C₆ alkyl, which is optionally substituted with one or more groups selected from R⁴ or Ht; C₂-C₆ alkenyl, which is optionally substituted with one or more groups selected from R⁴ or Ht; C₃-C₆ saturated carbocycle, which is optionally substituted with one or more groups selected from R⁴ or Ht; or C₅-C₆ unsaturated carbocycle, which is optionally substituted with one or more groups selected from R⁴ or Ht;

each R⁴ is independently selected from -R², -OR², -OR³, -SR², -SOR², -SO₂R², -CO₂R², -OC(O)-R², -C(O)-N(R²)₂, -C(O)-NR²(OR²), -S(O)₂-N(R²)₂, halo, -NR²-C(O)-R², -NR²-OR², -N(R²)₂ or -CN;

each R⁷ is independently selected from hydrogen,



wherein each M is independently selected from H, Li, Na, K, Mg, Ca, Ba, -N(R²)₄, C₁-C₁₂-alkyl, C₂-C₁₂-alkenyl, or -R⁶; wherein 1 to 4 -CH₂ radicals of the alkyl or alkenyl group, other than the -CH₂ that is bound to Z, is optionally replaced by a heteroatom group selected from O, S, S(O), S(O₂), or N(R²); and wherein any hydrogen in said alkyl, alkenyl or R⁶ is optionally replaced with a substituent selected from oxo, -C₁-C₄ alkyl, -N(R²)₂, -N(R²)₃, -OH, -O-(C₁-C₄ alkyl), -CN, -C(O)OR², -C(O)-N(R²)₂, S(O)₂-N(R²)₂, -N(R²)-C(O)-R₂, C(O)R², -S(O)_n-R², -OCF₃, -S(O)_n-R⁶, -N(R²)-S(O)₂(R²), halo, -CF₃, or -NO₂;

M' is H, C₁-C₁₂-alkyl, C₂-C₁₂-alkenyl, or -R⁶; wherein
1 to 4 -CH₂ radicals of the alkyl or alkenyl group is
optionally replaced by a heteroatom group selected from
O, S, S(O), S(O₂), or N(R²); and wherein any hydrogen in
5 said alkyl, alkenyl or R⁶ is optionally replaced with a
substituent selected from oxo, -OR², -C₁-C₄ alkyl, -N(R²)₂,
N(R²)₃, -OH, -O-(C₁-C₄ alkyl), -CN, -C(O)OR², -C(O)-N(R²)₂,
-S(O)₂-N(R²)₂, -N(R²)-C(O)-R₂, -C(O)R², -S(O)_n-R², -OCF₃,
-S(O)_n-R⁶, -N(R²)-S(O)₂(R²), halo, -CF₃, or -NO₂;

10 x is 0 or 1;

Z is O, S, N(R²)₂, or, when M is not present, H.

Y is P or S;

X is O or S; and

15 R⁹ is C(R²)₂, O or N(R²); and wherein when Y is S, Z
is not S; and

R⁶ is a 5-6 membered saturated, partially saturated
or unsaturated carbocyclic or heterocyclic ring system,
or an 8-10 membered saturated, partially saturated or
unsaturated bicyclic ring system; wherein any of said
20 heterocyclic ring systems contains one or more
heteroatoms selected from O, N, S, S(O)_n or N(R²); and
wherein any of said ring systems optionally contains 1 to
4 substituents independently selected from -OH, -C₁-C₄
alkyl, -O-(C₁-C₄ alkyl) or -O-C(O)-(C₁-C₄ alkyl).

25 It is also an object of this invention to provide
pharmaceutical compositions comprising the sulfonamides
of formula (I) and methods for their use as inhibitors of
HIV aspartyl protease.

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DETAILED DESCRIPTION OF THE INVENTION

In order that the invention herein described may be
more fully understood, the following detailed description

is set forth. In the description, the following terms are employed herein:

Unless expressly stated to the contrary, the terms "-SO₂-" and "-S(O)₂-" as used herein refer to a sulfone or sulfone derivative (i.e., both appended groups linked to the S), and not a sulfinate ester.

For the compounds of formula I, and intermediates thereof, the stereochemistry of OR⁷ is defined relative to D on the adjacent carbon atom, when the molecule is drawn in an extended zigzag representation (such as that drawn for compound of formula I). If both OR⁷ and D reside on the same side of the plane defined by the extended backbone of the compound, the stereochemistry of OR⁷ will be referred to as "syn". If OR⁷ and D reside on opposite sides of that plane, the stereochemistry of OR⁷ will be referred to as "anti".

The term "alkyl", alone or in combination with any other term, refers to a straight-chain or branch-chain saturated aliphatic hydrocarbon radical containing the specified number of carbon atoms, or where no number is specified, preferably from 1 to about 15 and more preferably from 1 to about 10 carbon atoms. Examples of alkyl radicals include, but are not limited to, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, pentyl, isoamyl, n-hexyl and the like.

The term "alkenyl," alone or in combination with any other term, refers to a straight-chain or branched-chain mono- or poly-unsaturated aliphatic hydrocarbon radical containing the specified number of carbon atoms, or where no number is specified, preferably from 2 to about 18 carbon atoms and more preferably, from 2 to about 8 carbon atoms. Examples of alkenyl radicals include, but

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are not limited to, ethenyl, propenyl, isopropenyl, 1,4-butadienyl, pentenyl and the like.

5 The term "alkoxy" refers to an alkyl ether radical, wherein the term "alkyl" is defined above. Examples of suitable alkyl ether radicals include, but are not limited to, methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, isobutoxy, sec-butoxy, tert-butoxy and the like.

10 The term "aryl," alone or in combination with any other term, refers to a carbocyclic aromatic radical (such as phenyl or naphthyl) containing the specified number of carbon atoms, preferably from 6-15 carbon atoms, and more preferably from 6-10 carbon atoms, optionally substituted with one or more substituents selected from C1-6 alkoxy, (for example methoxy), nitro, 15 cyano, -SCH₃, halogen, (for example chloro), amino, carboxylate and hydroxy. Examples of aryl radicals include, but are not limited to phenyl, naphthyl, indenyl, indanyl, azulenyl, fluorenyl, anthracenyl and the like.

20 The term "heterocyclyl" or "heterocycle" refers to a stable 3-7 membered monocyclic heterocyclic ring or 8-11 membered bicyclic heterocyclic ring which is either saturated or unsaturated, and which may be optionally benzofused if monocyclic. Each heterocycle consists of 25 one or more carbon atoms and from one to four heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur. As used herein, the terms "nitrogen and sulfur heteroatoms" include any oxidized form of nitrogen and sulfur, and the quaternized form of any basic 30 nitrogen. A heterocyclyl radical may be attached at any endocyclic carbon or heteroatom which results in the creation of a stable structure. Preferred heterocycles include 5-7 membered monocyclic heterocycles and 8-10

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membered bicyclic heterocycles. Examples of such groups include imidazolyl, imidazolinoyl, imidazolidinyl, quinolyl, isoquinolyl, indolyl, indazolyl, indazolinolyl, perhydropyridazyl, pyridazyl, pyridyl, pyrrolyl, 5 pyrrolinyl, pyrrolidinyl, pyrazolyl, pyrazinyl, quinoxolyl, piperidinyl, pyranyl, pyrazolinyl, piperazinyl, pyrimidinyl, pyridazinyl, morpholinyl, thiamorpholinyl, furyl, thienyl, triazolyl, thiazolyl, carbolinyl, tetrazolyl, thiazolidinyl, benzofuranoyl, 10 thiamorpholinyl sulfone, oxazolyl, benzoxazolyl, oxopiperidinyl, oxopyrrolidinyl, oxoazepinyl, azepinyl, isoxozolyl, isothiazolyl, furazanyl, tetrahydropyranyl, tetrahydrofuranlyl, thiazolyl, thiadiazoyl, dioxolyl, dioxinyl, oxathioly, benzodioxolyl, dithioly, 15 thiophenyl, tetrahydrothiophenyl, sulfolanyl, dioxanyl, dioxolanyl, tetrahydrofurodihydrofuranlyl, tetrahydropyranodihydrofuranlyl, dihydropyranyl, tetrahydrofurofuranlyl and tetrahydropyranofuranlyl.

The term "pharmaceutically effective amount" refers 20 to an amount effective in treating a virus infection, for example an HIV infection, in a patient either as monotherapy or in combination with other agents. The term "treating" as used herein refers to the alleviation of symptoms of a particular disorder in a patient or the 25 improvement of an ascertainable measurement associated with a particular disorder. The term "prophylactically effective amount" refers to an amount effective in preventing a virus infection, for example an HIV infection, in a patient. As used herein, the term 30 "patient" refers to a mammal, including a human.

The terms "HIV protease" and "HIV aspartyl protease" are used interchangeably and refer to the aspartyl protease encoded by the human immunodeficiency virus type

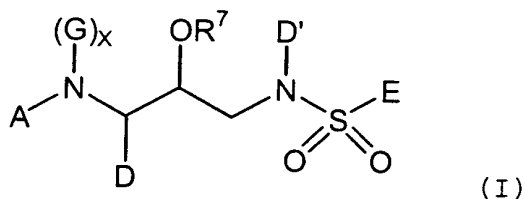
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1 or 2. In a preferred embodiment of this invention, these terms refer to the human immunodeficiency virus type 1 aspartyl protease.

The term "thiocarbamates" refers to compounds
5 containing the functional group N-SO₂-O.

Combinations of substituents and variables
envisioned by this invention are only those that result
in the formation of stable compounds. The term "stable",
as used herein, refers to compounds which possess
10 stability sufficient to allow manufacture and
administration to a mammal by methods known in the art.
Typically, such compounds are stable at a temperature of
40°C or less, in the absence of moisture or other
chemically reactive conditions, for at least a week.

15 This invention also envisions the quaternization of
any basic nitrogen-containing groups of the compounds
disclosed herein. The basic nitrogen can be quaternized
with any agents known to those of ordinary skill in the
art including, for example, lower alkyl halides, such as
20 methyl, ethyl, propyl and butyl chloride, bromides and
iodides; dialkyl sulfates including dimethyl, diethyl,
dibutyl and diamyl sulfates; long chain halides such as
decyl, lauryl, myristyl and stearyl chlorides, bromides
and iodides; and aralkyl halides including benzyl and
25 phenethyl bromides. Water or oil-soluble or dispersible
products may be obtained by such quaternization. The
novel sulfonamides of this invention are those of formula
I:



and pharmaceutically acceptable salts thereof;
wherein:

A is selected from H; Ht; $-R^1\text{-Ht}$; $-R^1\text{-C}_1\text{-C}_6$ alkyl, which is optionally substituted with one or more groups independently selected from hydroxy, $-\text{CN}$, $\text{C}_1\text{-C}_4$ alkoxy, Ht, $-\text{O-Ht}$, $-\text{NR}^2\text{-Ht}$, $-\text{NR}^2\text{-CO-N(R}^2\text{)}_2$, $-\text{SO}_2\text{-N(R}^2\text{)}_2$, $-\text{SO}_2\text{-R}^2$ or $-\text{CO-N(R}^2\text{)}_2$; $-R^1\text{-C}_2\text{-C}_6$ alkenyl, which is optionally substituted with one or more groups independently selected from hydroxy, $\text{C}_1\text{-C}_4$ alkoxy, Ht, $-\text{O-Ht}$, $-\text{NR}^2\text{-CO-N(R}^2\text{)}_2$ or $-\text{CO-N(R}^2\text{)}_2$; or R^7 ;

each R¹ is independently selected from -C(O)-, -S(O)₂-, -C(O)-C(O)-, -O-C(O)-, -O-S(O)₂-, -NR²-, -NR²-S(O)₂-, -NR²-C(O)- or -NR²-C(O)-C(O)-;

each Ht is independently selected from C₃-C₇ cycloalkyl; C₅-C₇ cycloalkenyl; C₆-C₁₄ aryl; or a 5-7 membered saturated or unsaturated heterocycle, containing one or more heteroatoms selected from N, N(R²), O, S and S(O)_n; wherein said aryl or said heterocycle is optionally fused to Q; and wherein any member of said Ht is optionally substituted with one or more substituents independently selected from oxo, -OR², SR², -R², -N(R²)(R²), -R²-OH, -CN, -CO₂R², -C(O)-N(R²)₂, -S(O)₂-N(R²)₂, -N(R²)-C(O)-R², -N(R²)-C(O)O-R², -C(O)-R², -S(O)_n-R², -OCF₃, -S(O)_n-Q, methylenedioxy, -N(R²)-S(O)₂(R²), halo, -CF₃, -NO₂, Q, -OQ, -OR⁷, -SR⁷, -R⁷, -N(R²)(R⁷) or -N(R⁷)₂;

each R² is independently selected from H, or C₁-C₄ alkyl optionally substituted with a 3-7 membered saturated, partially saturated or unsaturated carbocyclic ring system; or a 5-7 membered saturated, partially saturated or unsaturated heterocyclic ring containing one or more heteroatoms selected from O, N, S, S(O)_n or N(R³³); wherein any of said ring systems or N(R³³) is optionally substituted with 1 to 4 substituents

independently selected from -X'-Y', -O-arylalkyl,
-S-arylalkyl, -N(Y')₂, -N(H)-arylalkyl, -N(C₁-C₄
alkyl)-arylalkyl, oxo, -O-(C₁-C₄ alkyl), OH, C₁-C₄ alkyl,
-SO₂H, -SO₂-(C₁-C₄ alkyl), -SO₂-NH₂, -SO₂-NH(C₁-C₄ alkyl),
5 -SO₂-N(C₁-C₄ alkyl)₂, -NH₂, -NH(C₁-C₄ alkyl), -N(C₁-C₄
alkyl)₂, -NH-C(O)H, -N(C₁-C₄ alkyl)-C(O)H, -NH-C(O)-C₁-C₄
alkyl, -C₁-C₄ alkyl-OH, -OH, -CN, -C(O)OH, -C(O)O-C₁-C₄
alkyl, -C(O)-NH₂, -C(O)-NH(C₁-C₄ alkyl), -C(O)-N(C₁-C₄
alkyl)₂, halo or -CF₃;

10 X' is -O-, -S-, -NH-, -NHC(O)-, -NHC(O)O-, -NHSO₂-,
or -N(C₁-C₄)alkyl-;

Y' is C₁-C₁₅ alkyl, C₂-C₁₅ alkenyl or alkynyl, wherein
one to five carbon atoms in Y are optionally substituted
with C₃-C₇ cycloalkyl or C₅-C₆ cycloalkenyl, C₆-C₁₄ aryl or
15 a 5-7 membered saturated or unsaturated heterocycle,
containing one or more heteroatoms selected from N, NH,
O, S and S(O)_n;

each R³ is independently selected from H, Ht, C₁-C₆
alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₆ cycloalkyl or
20 C₅-C₆ cycloalkenyl; wherein any member of said R³, except
H, is optionally substituted with one or more
substituents selected from -OR², -C(O)-N(R²)₂,
-S(O)_n-N(R²)₂, -N(R²)₂, -N(R²)-C(O)O(R²), -N(R²)-C(O)N(R²)₂,
-N(R²)-C(O)-R², Ht, -CN, -SR², -C(O)OR², N(R²)-C(O)-R²;

25 each R³³ is selected from H, C₁-C₆ alkyl, C₂-C₆
alkenyl, C₂-C₆ alkynyl, C₃-C₆ cycloalkyl or C₅-C₆
cycloalkenyl, C₆-C₁₄ aryl or a 5-7 membered saturated or
unsaturated heterocycle, containing one or more
heteroatoms selected from N, NH, O, S and S(O)_n;

30 each n is independently 1 or 2;

G, when present, is selected from H, R⁷ or C₁-C₄
alkyl, or, when G is C₁-C₄ alkyl, G and R⁷ are bound to one

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another either directly or through a C₁-C₃ linker to form a heterocyclic ring; or

when G is not present (i.e., when x in (G)_x is 0), then the nitrogen to which G is attached is bound
5 directly to the R⁷ group in -OR⁷ with the concomitant displacement of one -ZM group from R⁷;

D is selected from C₁-C₆ alkyl which is substituted with Q, which is optionally substituted with one or more groups selected from C₃-C₆ cycloalkyl, -R³, -O-Q or Q;
10 C₂-C₄ alkenyl which is substituted with Q, which is optionally substituted with one or more groups selected from -OR², -S-Ht, -R³, -O-Q or Q; C₃-C₆ cycloalkyl, which is optionally substituted with or fused to Q; or C₅-C₆ cycloalkenyl, which is optionally substituted with or
15 fused to Q;

each Q is independently selected from a 3-7 membered saturated, partially saturated or unsaturated carbocyclic ring system; or a 5-7 membered saturated, partially
20 saturated or unsaturated heterocyclic ring containing one or more heteroatoms selected from O, N, S, S(O)_n or N(R²); wherein Q contains one substituent selected from -OR², -OR⁸, -O-arylalkyl, -SR⁸, -S-arylalkyl, -N(R²)R⁸, -N(R²)-arylalkyl and may be optionally substituted with one or more additional substituents independently selected
25 from oxo, -OR⁸, -O-arylalkyl -SR⁸, -S-arylalkyl, -N(R²)R⁸, -N(R²)-arylalkyl, -OR², -R², -SO₂R², -SO₂-N(R²)₂, -N(R²)₂, -N(R²)-C(O)-R², -OH, (C₁-C₄)-OH, -CN, -CO₂R², -C(O)-N(R²)₂, halo or -CF₃;

each R⁸ is independently selected from Ht, -C₁-C₁₅
30 branched or straight chain alkyl, alkenyl or alkynyl wherein one to five carbon atoms in said alkyl, alkenyl or alkynyl are independently replaced by W, or wherein one to five carbon atoms in said alkyl, alkenyl or

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alkynyl are substituted with Ht; and wherein R⁸ is additionally and optionally substituted with one or more groups independently selected from -OH, -S(C₁-C₆ alkyl), -CN, -CF₃, -N(R²)₂, halo, -C₁-C₄-alkyl, -C₁-C₄-alkoxy; -Ht; -O-Ht; -NR²-CO-N(R²)₂; -CO-N(R²)₂; -R¹-C₂-C₆ alkenyl, which is optionally substituted with one or more groups independently selected from hydroxy, C₁-C₄ alkoxy, Ht, -O-Ht, -NR²-CO-N(R²)₂ or -CO-N(R²)₂; or R⁷;

wherein W is -O-, -NR²-, -S-, -C(O)-, -C(S)-, -C(=NR²)-, -S(O)₂-, -NR²-S(O)₂-, -S(O)₂-NR²-, -NR²-C(O)O-, -O-C(O)NR²-, -NR²-C(O)NR²-, -NR²-C(S)NR²-, -CONR², -NR²C(O)-, -C(S)NR², -NR²C(S)-, -NR²-C(=N-CN)-NR²-, -NR²C(=N-CN)O- or -C(O)O-;

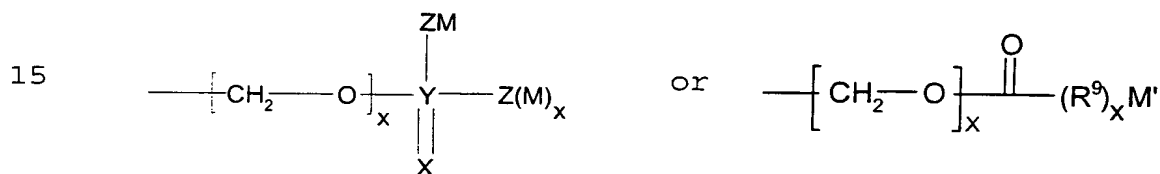
D' is selected from C₁-C₁₅ alkyl, C₁-C₁₅ alkoxy, C₂-C₁₅ alkenyl, C₂-C₁₅ alkenyloxy, C₂-C₁₅ alkynyl, or C₂-C₁₅ alkynyloxy, wherein D' optionally comprises one or more substituents independently selected from Ht, oxo, halo, -CF₃, -OCF₃, -NO₂, azido, -SH, -SR³, -N(R³)-N(R³)₂, -O-N(R³)₂, -(R³)N-O-(R³), -N(R³)₂, -CN, -CO₂R³, -C(O)-N(R³)₂, -S(O)_n-N(R³)₂, -N(R³)-C(O)-R³, -N(R³)-C(O)-N(R³)₂, -C(O)-R³, -S(O)_n-R³, -N(R³)-S(O)_n(R³), -N(R³)-S(O)_n-N(R³)₂, -S-NR³-C(O)R³, -C(S)N(R³)₂, -C(S)R³, -NR³-C(O)OR³, -O-C(O)OR³, -O-C(O)N(R³)₂, -NR³-C(S)R³, =N-OH, =N-OR³, =N-N(R³)₂, =NR³, =NNR³C(O)N(R³)₂, =NNR³C(O)OR³, =NNR³S(O)_n-N(R³)₂, -NR³-C(S)OR³, -NR³-C(S)N(R³)₂, -NR³-C[=N(R³)]-N(R³)₂, -N(R³)-C[=N-NO₂]-N(R³)₂, -N(R³)-C[=N-NO₂]-OR³, -OC(O)R³, -OC(S)R³, -OC(O)N(R³)₂, -C(O)N(R³)-N(R³)₂, -N(R³)-N(R³)C(O)R³, -N(R³)-OC(O)R³, -N(R³)-OC(O)R³, -N(R³)-OC(S)N(R³)₂, -OC(S)N(R³)(R³), or -PO₃-R³;

E is selected from Ht; O-Ht; Ht-Ht; Ht fused with Ht; -O-R³; -N(R²)(R³); -N(R²)-Ht; C₁-C₆ alkyl, which is optionally substituted with one or more groups selected

from R⁴ or Ht; C₂-C₆ alkenyl, which is optionally substituted with one or more groups selected from R⁴ or Ht; C₃-C₆ saturated carbocycle, which is optionally substituted with one or more groups selected from R⁴ or Ht; or C₅-C₆ unsaturated carbocycle, which is optionally substituted with one or more groups selected from R⁴ or Ht;

each R⁴ is independently selected from -R², -OR², -OR³, -SR², -SOR², -SO₂R², -CO₂R², -OC(O)-R², -C(O)-N(R²)₂, -C(O)-NR²(OR²), -S(O)₂-N(R²)₂, halo, -NR²-C(O)-R², -NR²-OR², -N(R²)₂ or -CN;

each R⁷ is independently selected from hydrogen,



wherein each M is independently selected from H, Li, Na, K, Mg, Ca, Ba, -N(R²)₄, C₁-C₁₂-alkyl, C₂-C₁₂-alkenyl, or -R⁶; wherein 1 to 4 -CH₂ radicals of the alkyl or alkenyl group, other than the -CH₂ that is bound to Z, is optionally replaced by a heteroatom group selected from O, S, S(O), S(O₂), or N(R²); and wherein any hydrogen in said alkyl, alkenyl or R⁶ is optionally replaced with a substituent selected from oxo, -C₁-C₄ alkyl, -N(R²)₂, -N(R²)₃, -OH, -O-(C₁-C₄ alkyl), -CN, -C(O)OR², -C(O)-N(R²)₂, S(O)₂-N(R²)₂, -N(R²)-C(O)-R₂, C(O)R², -S(O)_n-R², -OCF₃, -S(O)_n-R⁶, -N(R²)-S(O)₂(R²), halo, -CF₃, or -NO₂;

M' is H, C₁-C₁₂-alkyl, C₂-C₁₂-alkenyl, or -R⁶; wherein 1 to 4 -CH₂ radicals of the alkyl or alkenyl group is optionally replaced by a heteroatom group selected from

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O, S, S(O), S(O₂), or N(R²); and wherein any hydrogen in said alkyl, alkenyl or R⁶ is optionally replaced with a substituent selected from oxo, -OR², -C₁-C₄ alkyl, -N(R²)₂, N(R²)₃, -OH, -O-(C₁-C₄ alkyl), -CN, -C(O)OR², -C(O)-N(R²)₂,
 5 -S(O)₂-N(R²)₂, -N(R²)-C(O)-R₂, -C(O)R², -S(O)_n-R², -OCF₃,
 -S(O)_n-R⁶, -N(R²)-S(O)₂(R²), halo, -CF₃, or -NO₂;

x is 0 or 1;

Z is O, S, N(R²)₂, or, when M is not present, H.

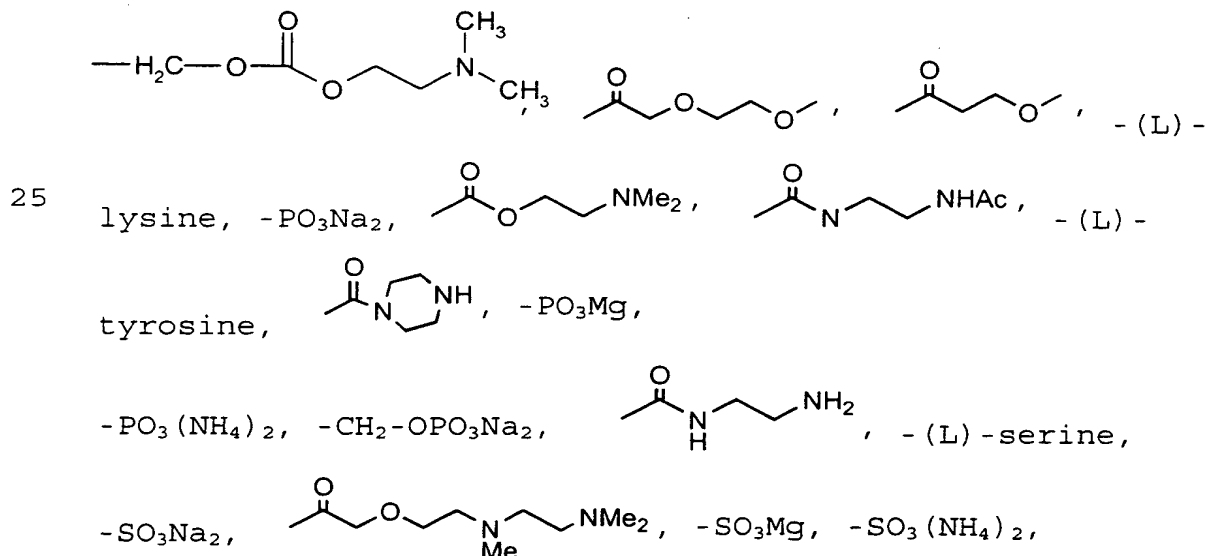
Y is P or S;

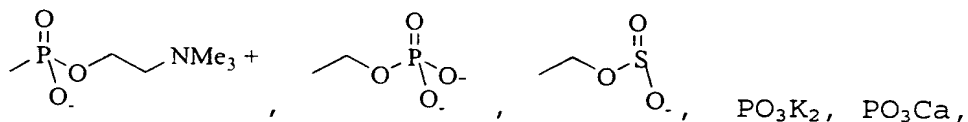
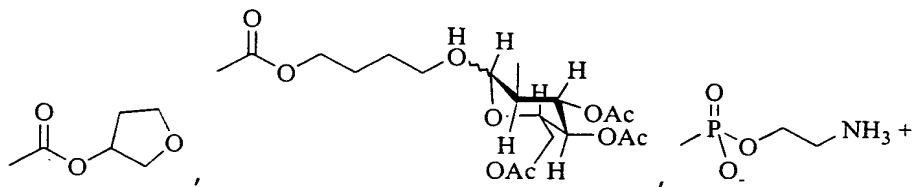
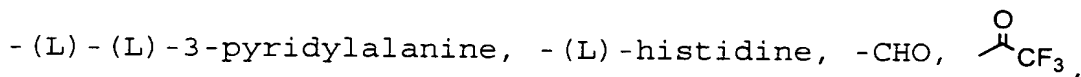
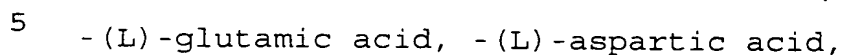
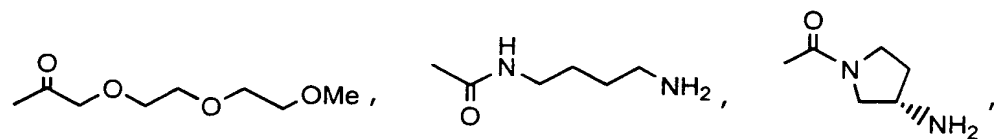
10 X is O or S; and

R⁹ is C(R²)₂, O or N(R²); and wherein when Y is S, Z is not S; and

R⁶ is a 5-6 membered saturated, partially saturated or unsaturated carbocyclic or heterocyclic ring system,
 15 or an 8-10 membered saturated, partially saturated or unsaturated bicyclic ring system; wherein any of said heterocyclic ring systems contains one or more heteroatoms selected from O, N, S, S(O)_n or N(R²); and wherein any of said ring systems optionally contains 1 to
 20 4 substituents independently selected from -OH, -C₁-C₄ alkyl, -O-(C₁-C₄ alkyl) or -O-C(O)-(C₁-C₄ alkyl).

Preferably, at least one R⁷ is selected from:

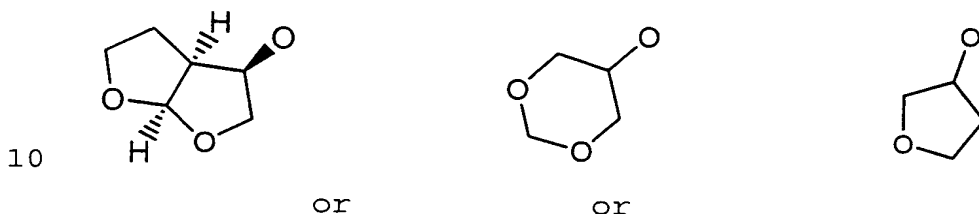




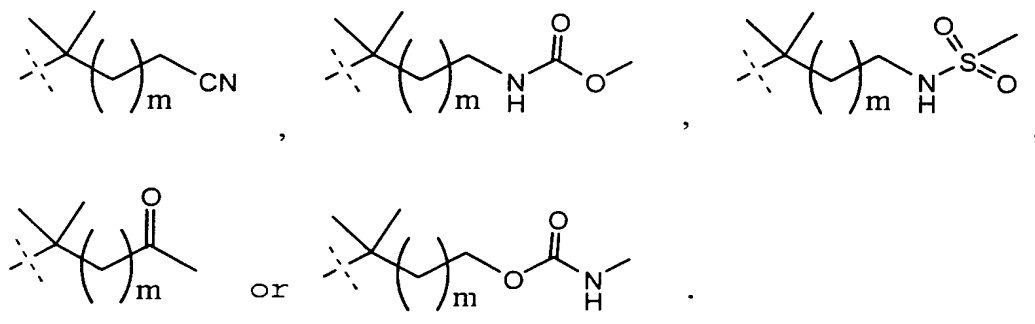
It will be understood by those of skill in the art that component M or M' in the formulae set forth herein will have either a covalent, a covalent/ zwitterionic, or an ionic association with either Z or R⁹ depending upon the actual choice for M or M'. When M or M' is hydrogen, alkyl, alkenyl, or R⁶, M or M' is covalently bound to R⁹ or Z. If M is a mono- or bivalent metal or other charged species (i.e., NH₄⁺), there is an ionic interaction between M and Z and the resulting compound is a salt.

When x is 0 in $(M)_x$, Z may be a charged species.
 When that occurs, the other M may be oppositely charged
 to produce a 0 net charge on the molecule.
 Alternatively, the counter ion may located elsewhere in
 5 the molecule.

According to yet another preferred embodiment, A is
 $R'-C(O)$, wherein R' is selected from any of the R' groups
 indicated in Tables 1, 2 and 3, below. More preferably,
 R' is selected from $-R^1-C_1-C_6$ alkyl,



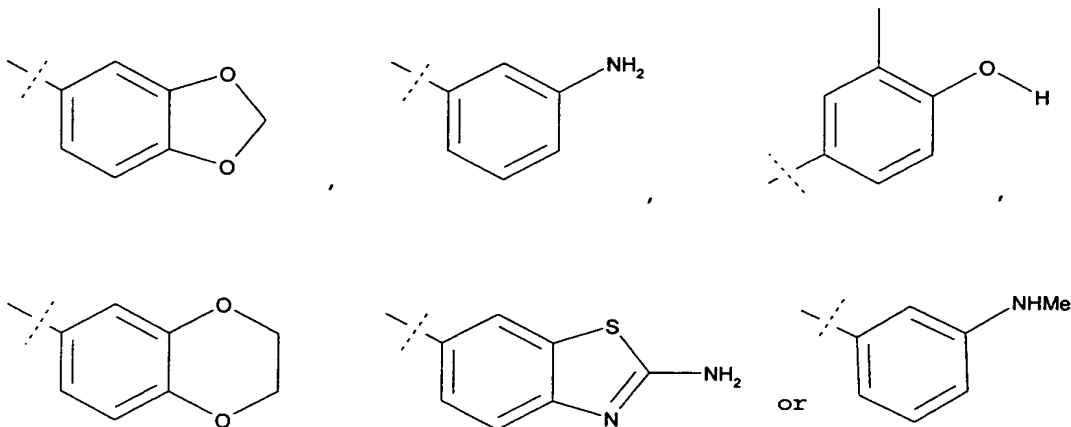
In another preferred embodiment, D' is $-CH_2-R''$,
 wherein R'' is selected from any of the R'' groups
 indicated in Tables 1, 2 and 3, below. More preferably,
 15 R'' is selected from:



wherein m is 0 to 3.

According to another preferred embodiment, E is
 selected from any of the E groups indicated in Tables 1,

2 and 3, below. More preferably, E is selected from:



Preferably, W is $-O-$, $-NR^2-$, $-NR^2-S(O)_2-$, $-NR^2-C(O)O-$,
 $-O-C(O)NR^2-$, $-NR^2-C(O)NR^2-$, $-NR^2-C(S)NR^2-$, $-NR^2C(O)-$,
 5 $-C(O)NR^2$, $-NR^2-C(=N-CN)-NR^2-$, $-NR^2C(=N-CN)O-$ or $-C(O)O-$.

More preferably, W is $-NR^2-$, $-NR^2C(O)-$ or $-C(O)NR^2$.
 Most preferably, W is $-NH-$, $-NHC(O)$ or $-C(O)NH-$.

According to a preferred embodiment, R^8 is $-C_1-C_4-$
 branched or straight chain alkyl, wherein one to two
 10 carbon atoms in said alkyl are independently replaced by
 W, wherein R^8 is additionally and optionally substituted
 with one or more groups independently selected from $-OH$;
 $-C_1-C_4$ -alkoxy; $-Ht$; $-O-Ht$; $-NR^2-CO-N(R^2)_2$; $-CO-N(R^2)_2$;
 $-R^1-C_2-C_6$ alkenyl, which is optionally substituted with
 15 one or more groups independently selected from hydroxy,
 C_1-C_4 alkoxy, Ht , $-O-Ht$, $-NR^2-CO-N(R^2)_2$; $-CO-N(R^2)_2$; or R^7 .

According to another preferred embodiment, R^8 is Ht ,
 wherein Ht is C_{6-14} aryl or a 5-7 membered saturated or
 unsaturated heterocycle, containing one or more
 20 heteroatoms selected from N , $N(R^2)$, O , S and $S(O)_n$,
 wherein Ht is optionally substituted as defined above.

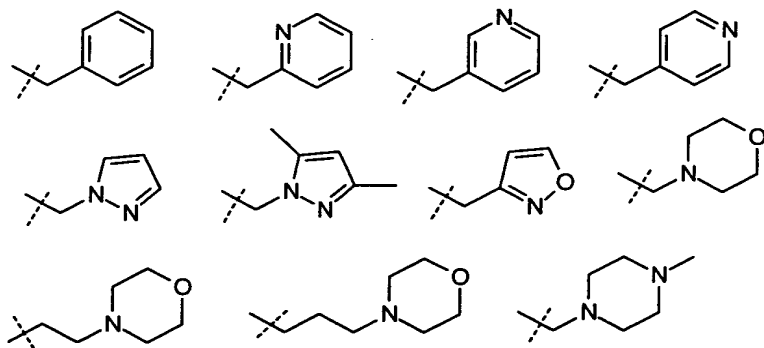
According to another preferred embodiment, R^8 is a
 $-C_1-C_4$ -branched or straight alkyl chain, wherein one to
 two carbon atoms are substituted with Ht , wherein Ht is
 25 C_{6-14} aryl or a 5-7 membered saturated or unsaturated

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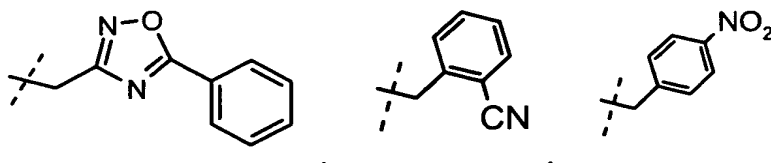
heterocycle, containing one or more heteroatoms selected from N, N(R²), O, S and S(O)_n, wherein Ht is optionally substituted as defined above.

According to a more preferred embodiment, R⁸ is a
 5 -C₁-C₄-branched or straight alkyl chain, wherein one carbon atom in said alkyl chain is substituted with Ht, wherein Ht is phenyl or Ht is a 5-6 membered saturated or unsaturated heterocycle, containing one or more heteroatoms selected from N, N(R²), O and S, wherein Ht is
 10 optionally substituted as defined above. Preferably, said Ht in R⁸ is selected from 2-thienyl, 3-thienyl, 2-furyl, 3-furyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, thiazolyl, morpholinyl, pyrimidinyl, thiazolidinyl, imidazolyl, 1,2,3-triazolyl, pyrrolidinyl, pyrazolyl,
 15 piperazyl, 1,2,4-oxadiazolyl, 4-4'-thiadiazolyl, 1,2,3-thiadiazolyl, isoxazolyl, and isothiazolyl, wherein said Ht is optionally substituted, as defined above. More preferably, R⁸ is selected from any of the R⁸ groups depicted in Tables 1, 2 and 3.

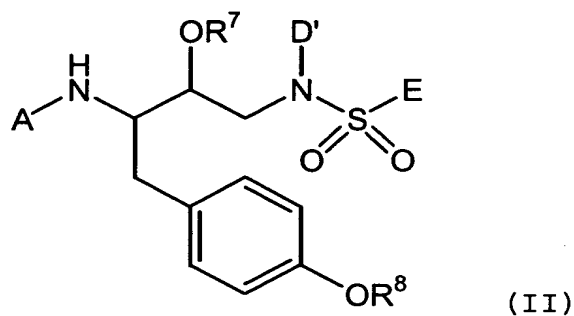
20 Most preferably, R⁸ is selected from:







More preferred compounds of formula I are those represented by formula II:



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wherein A, R⁷, D', E and R⁸ are as defined above.

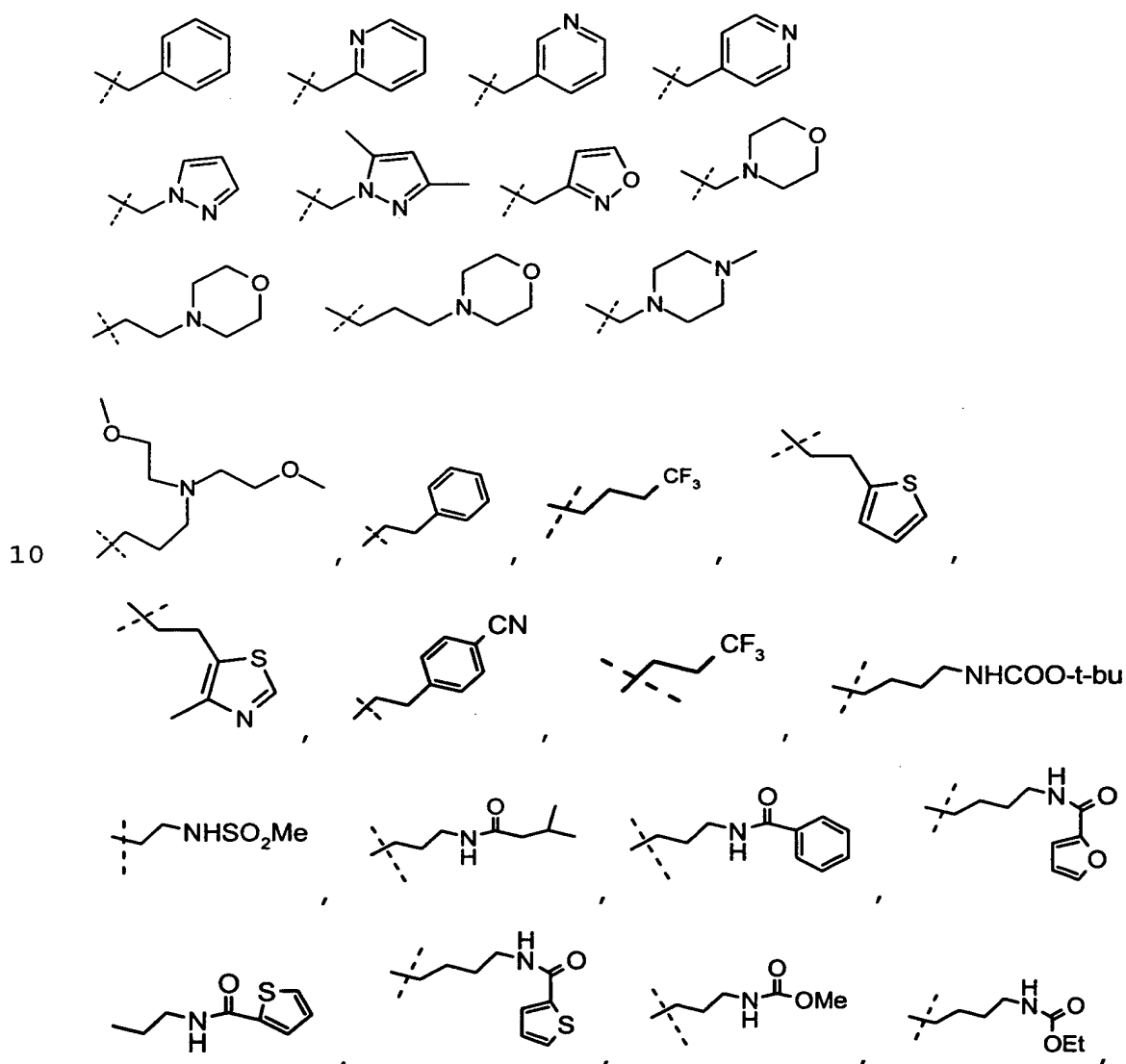
Preferred compounds of formula II set forth above are those, wherein R⁷ in -OR⁷ is -PO(OM)₂ or

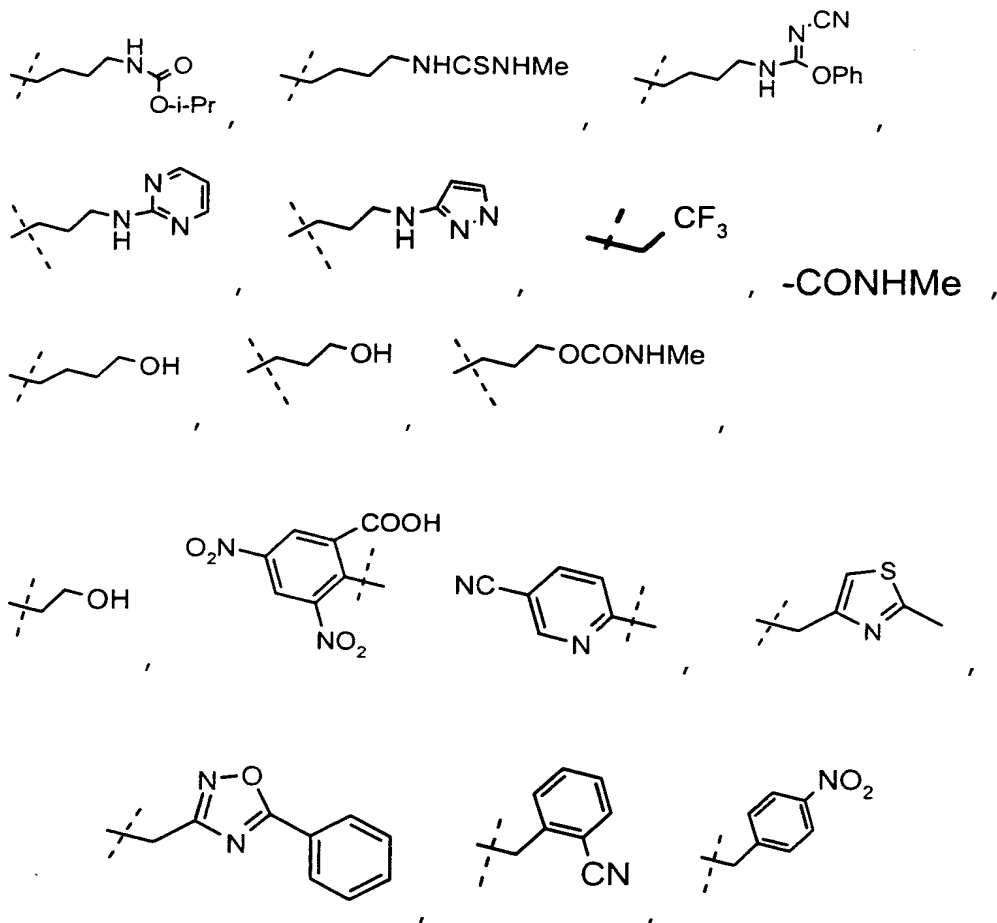
10 C(O)CH₂OCH₂CH₂OCH₂CH₂OCH₃; wherein M is H, Li, Na, Ca, Mg, K or C₁-C₄ alkyl.

Also preferred are compounds of formula II, wherein R⁸ is selected from -C₁-C₁₅ branched or straight chain alkyl, alkenyl or alkynyl wherein one to five carbon atoms in said alkyl, alkenyl or alkynyl are independently replaced by W, or wherein one to five carbon atoms in said alkyl, alkenyl or alkynyl are substituted with Ht; and wherein R⁸ is additionally and optionally substituted with one or more groups independently selected from -OH, 15 -SCH₃, -CN, -CF₃, amino, halo, -C₁-C₄-alkyl, -C₁-C₄-alkoxy; -Ht; -O-Ht; -NR²-CO-N(R²)₂; -CO-N(R²)₂; -R¹-C₂-C₆ alkenyl, 20 which is optionally substituted with one or more groups

More preferably, R⁸ in compounds of formula II is selected from any of the R⁸ groups depicted in Tables 1, 2 and 3. According to another more preferred embodiment, R⁸ in compounds of formula II is selected from the group consisting of:

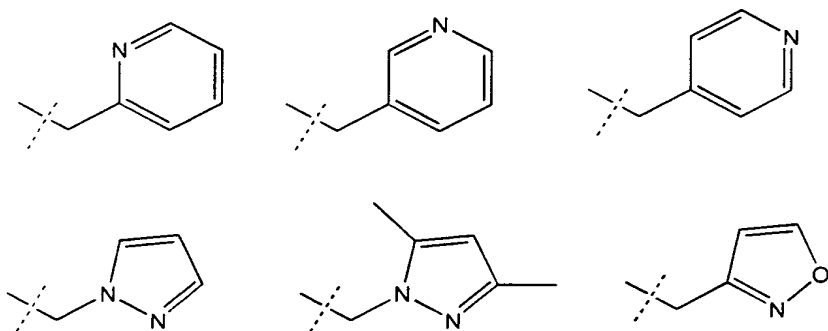
Most preferably, R⁸ is selected from:





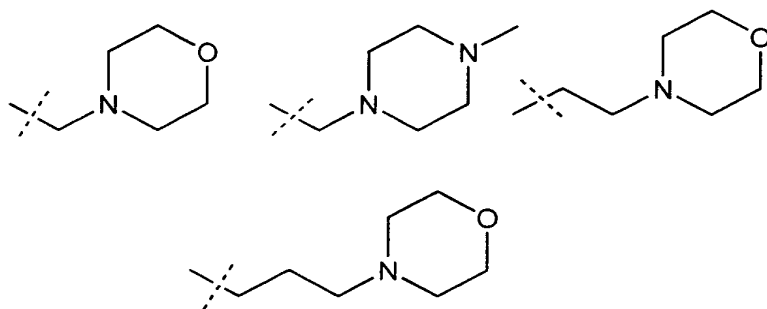
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According to another more preferred embodiment, R⁸ is
 10 selected from:





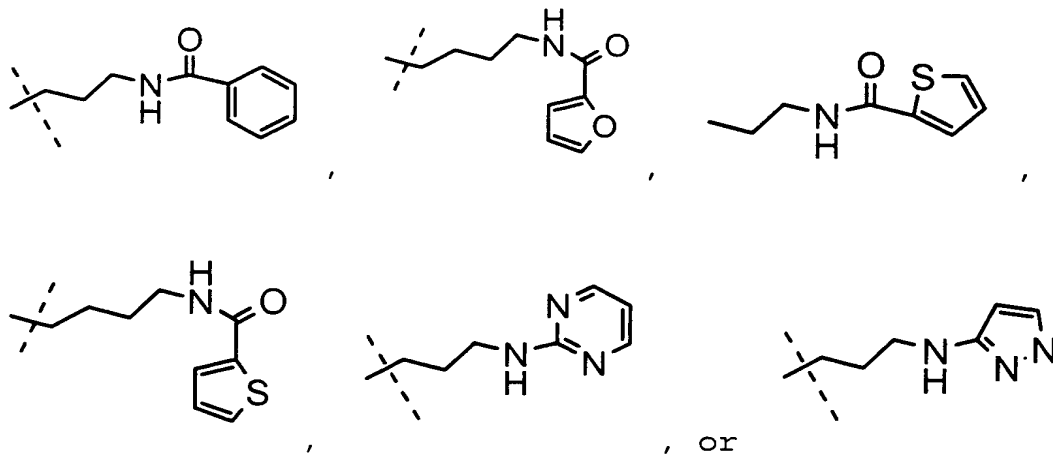
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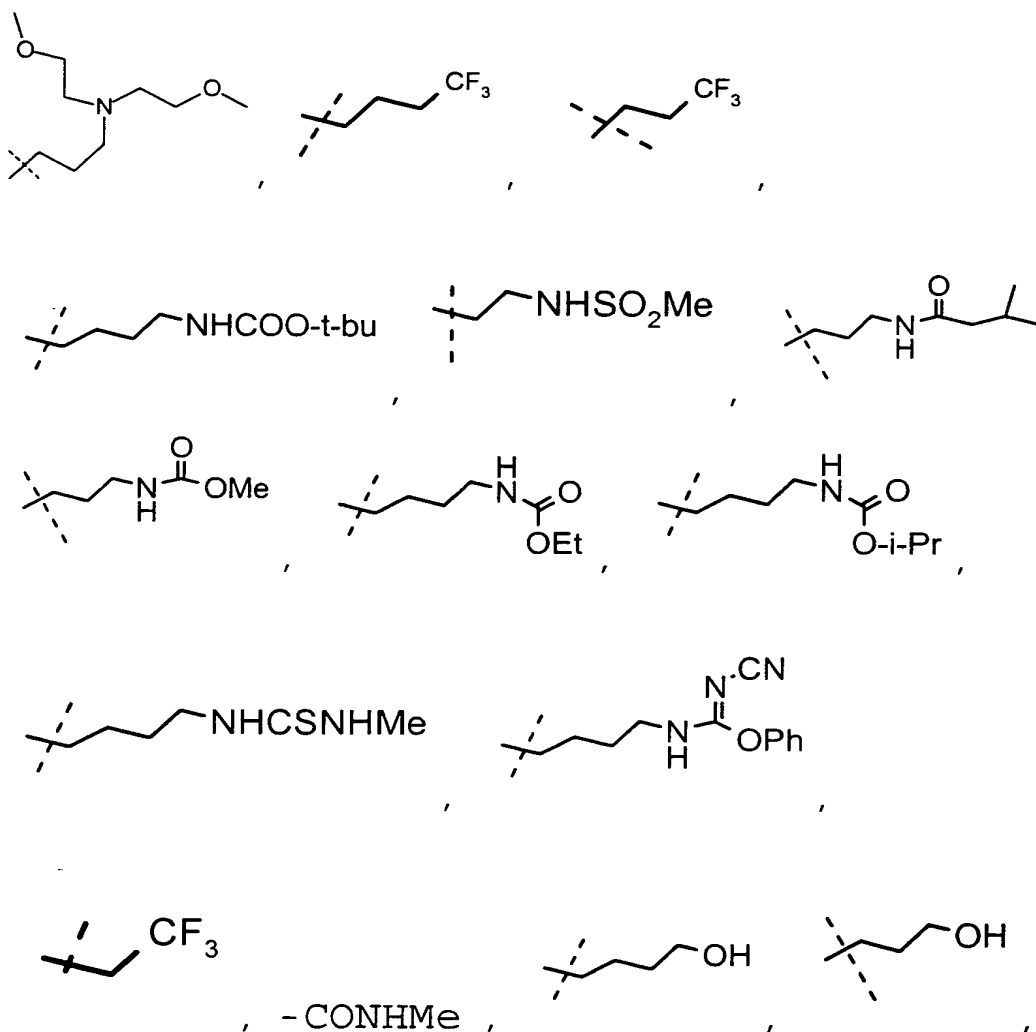
selected from:

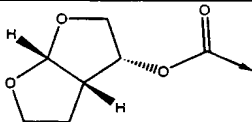
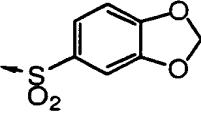
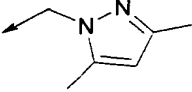
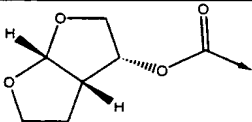
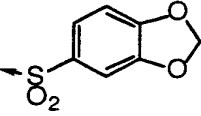
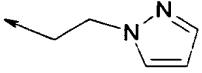
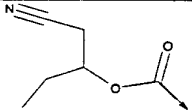
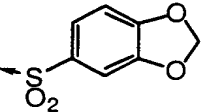
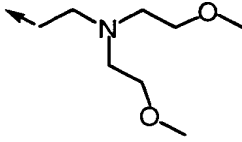
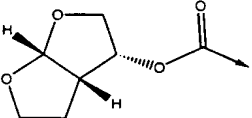
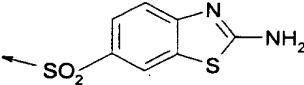
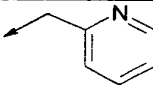
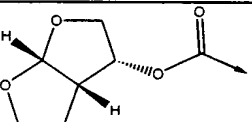
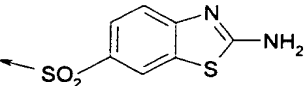
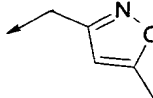
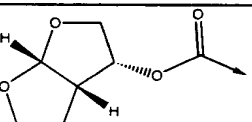
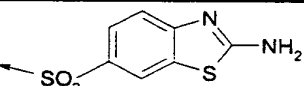
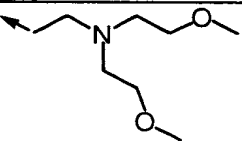
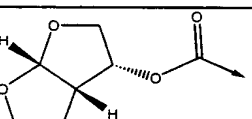
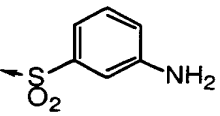
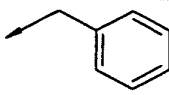
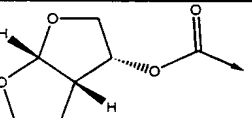
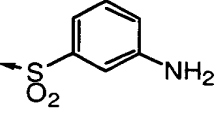
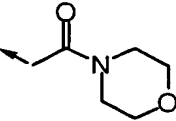


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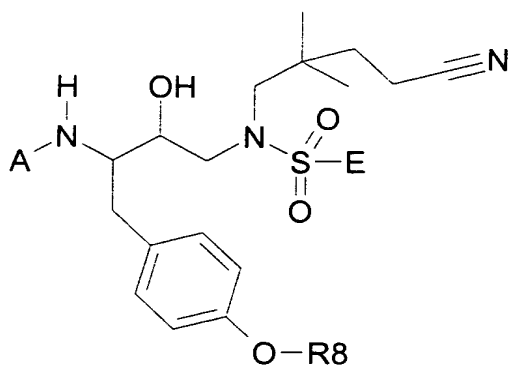
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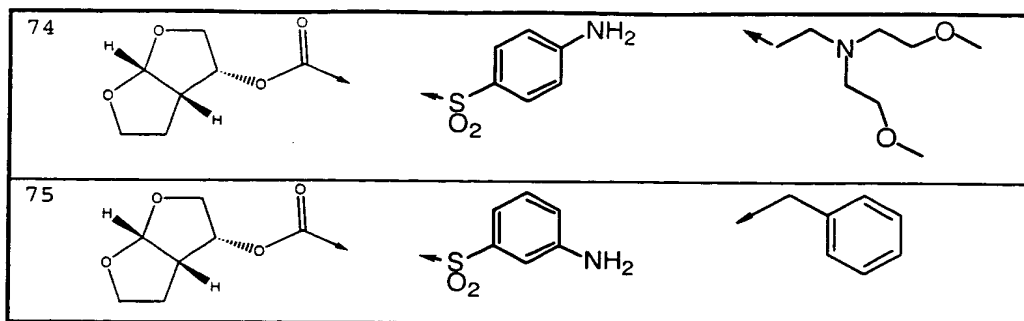
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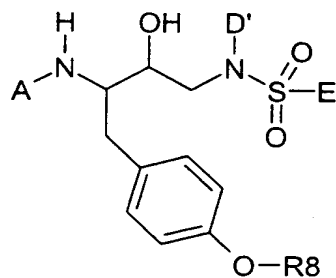


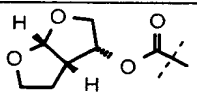

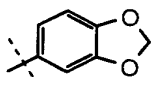
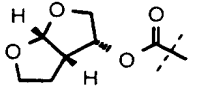
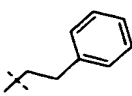

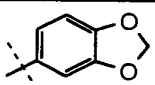
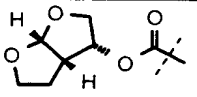
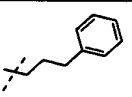

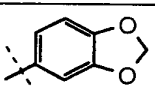
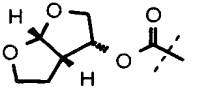
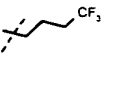

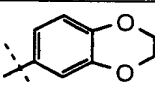
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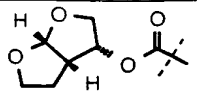
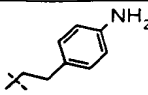

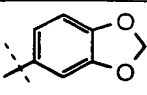
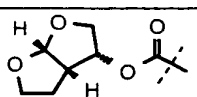
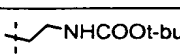

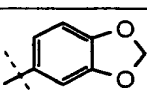
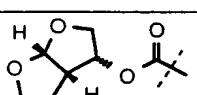


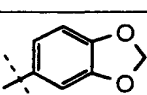
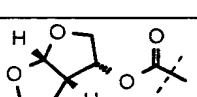
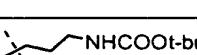

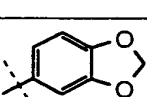
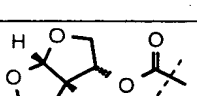
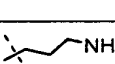

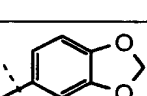
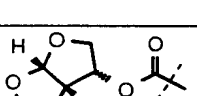
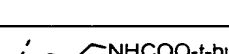

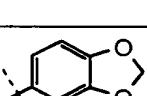
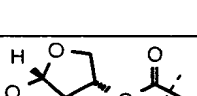
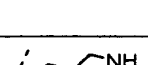

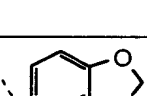
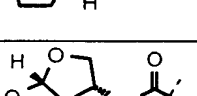
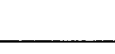

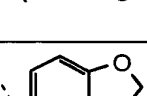
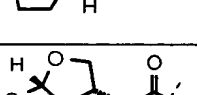
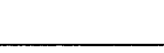

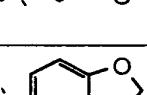
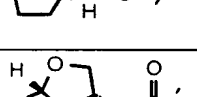


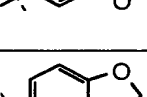
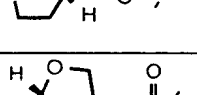


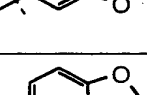
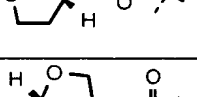


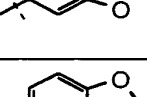
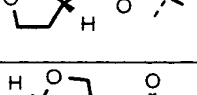


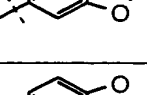
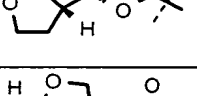
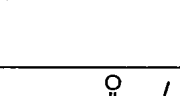

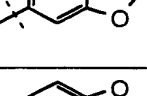
5 Table 3

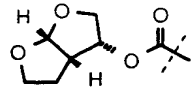
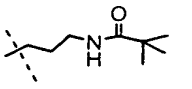

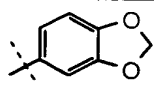
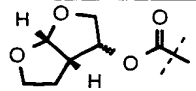
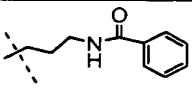

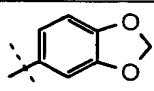
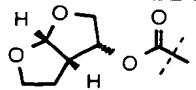
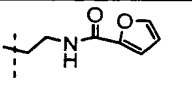

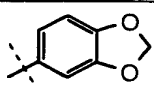
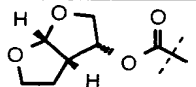
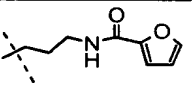

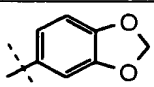
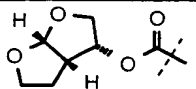
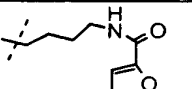

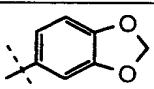
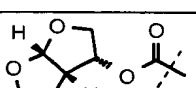
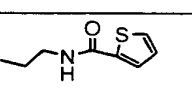

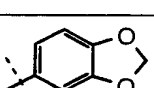
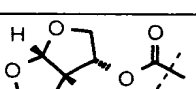
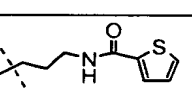

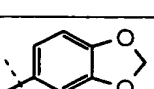
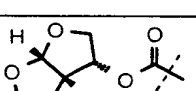
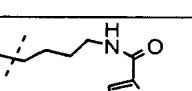

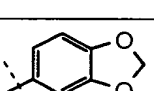
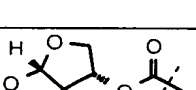
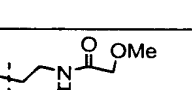

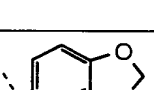
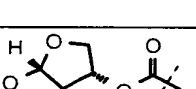
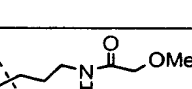

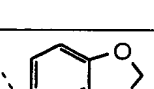
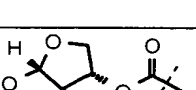
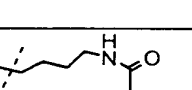

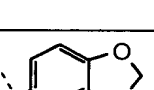
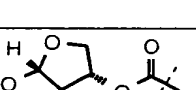
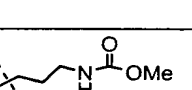

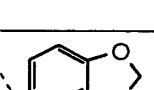
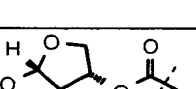
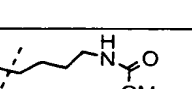

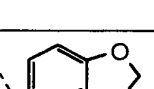


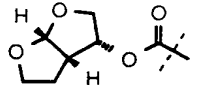
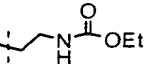

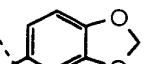
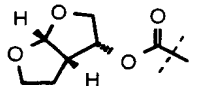
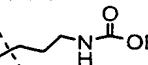

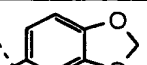
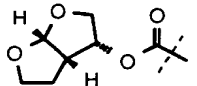
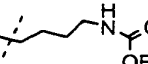

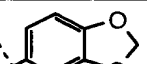
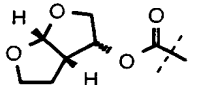
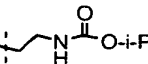

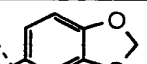
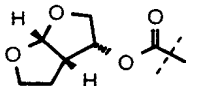
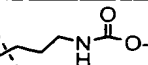

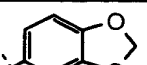
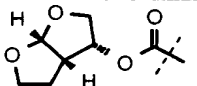
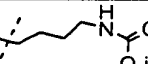


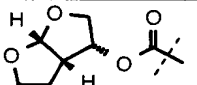
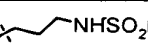

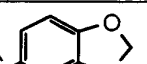
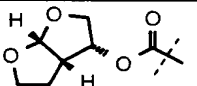
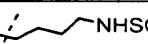

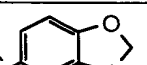
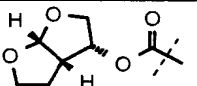


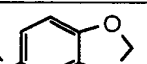
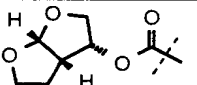
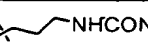

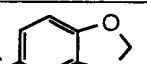
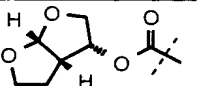
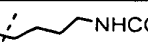

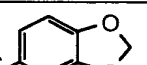
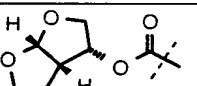
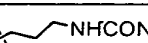

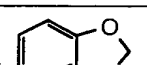
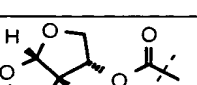
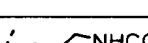

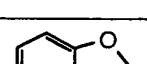
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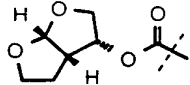


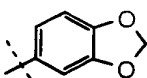
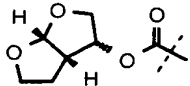
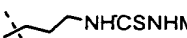

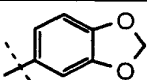
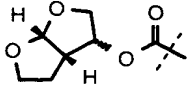
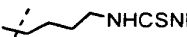

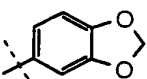
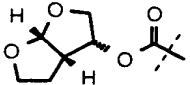
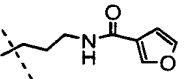

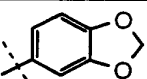
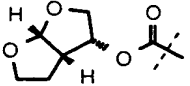
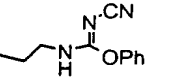

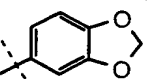
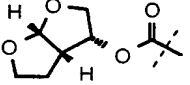
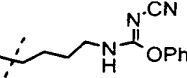

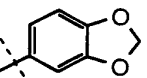
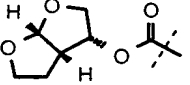
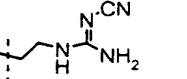

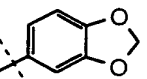
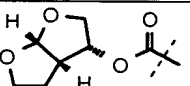
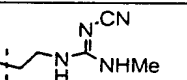

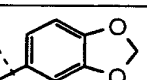
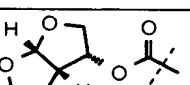
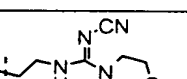

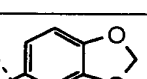
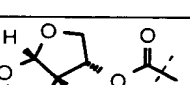
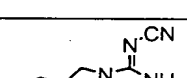

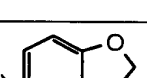
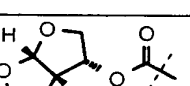
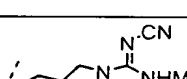

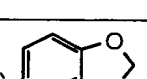
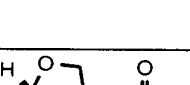
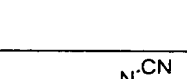

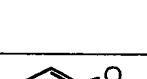
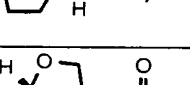
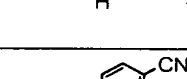

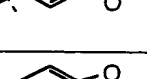
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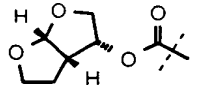
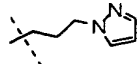

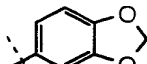
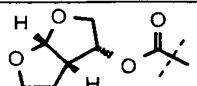
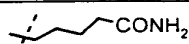

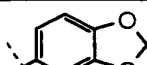
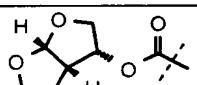
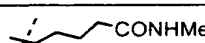

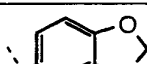
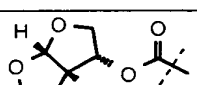
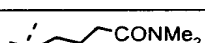

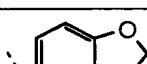
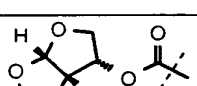
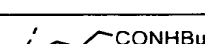

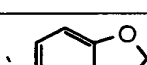
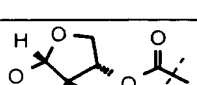
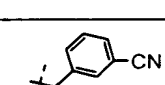

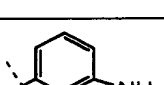
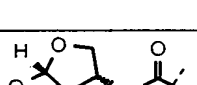
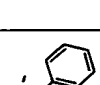


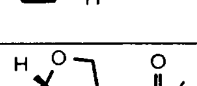

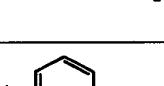
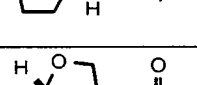


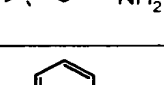
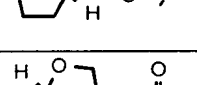
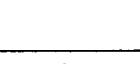

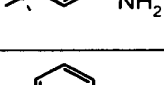
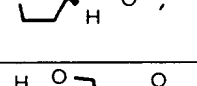


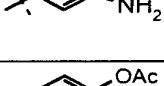
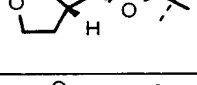
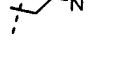

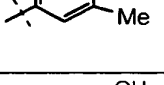
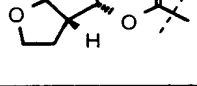
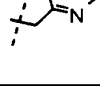

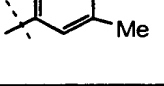
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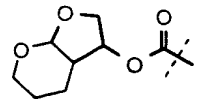


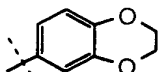
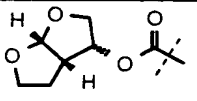

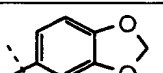
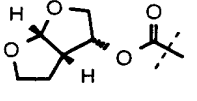

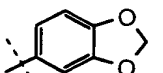
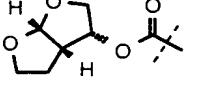

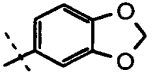
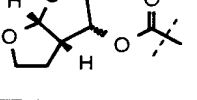

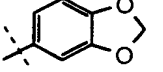
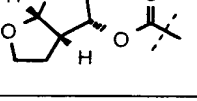
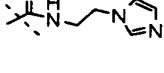

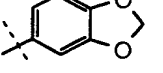
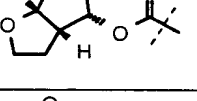
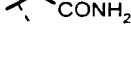

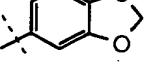
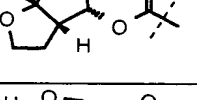
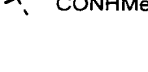

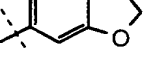
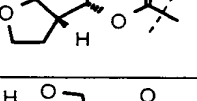


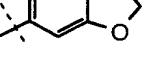
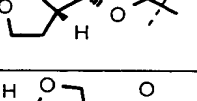


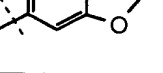
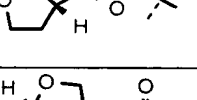
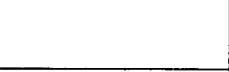

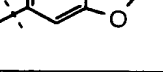
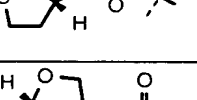


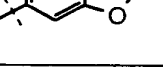
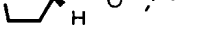


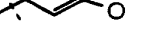
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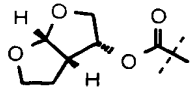
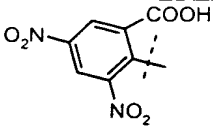
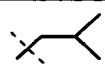
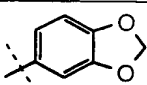
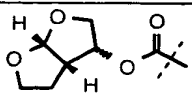
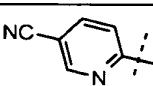
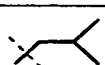
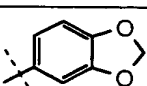
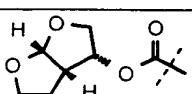
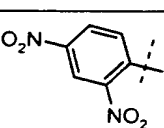
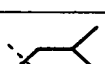
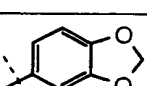
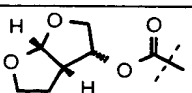
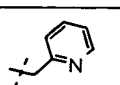
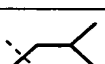
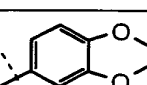
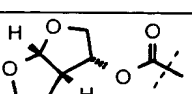
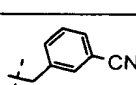
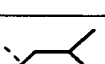
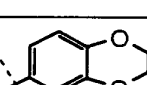
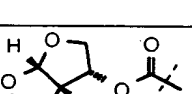
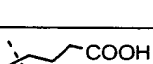

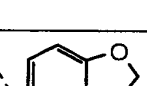
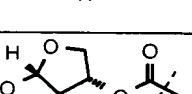

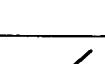
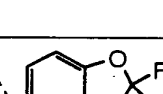
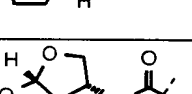

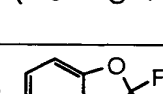
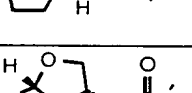
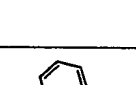

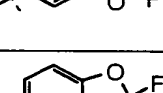
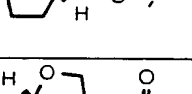


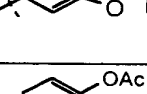
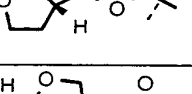

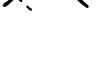
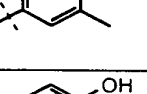
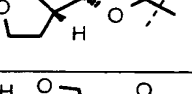
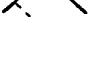
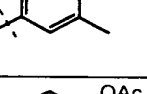
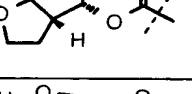
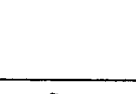
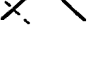
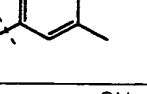
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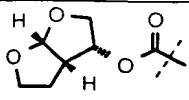

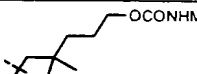
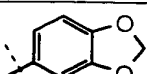
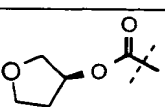

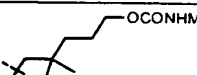
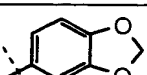
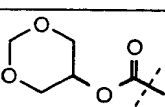

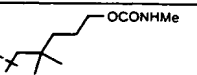
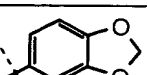
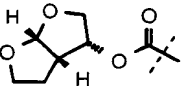


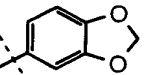
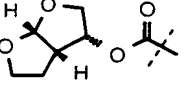


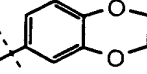
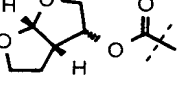
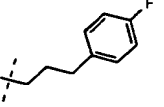

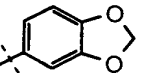
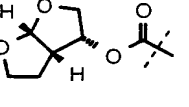
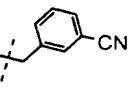

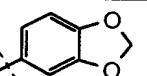
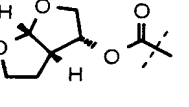
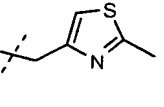

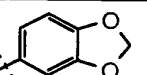
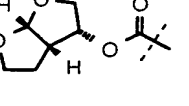
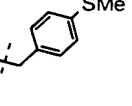

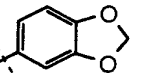
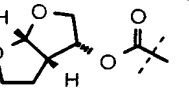
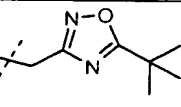

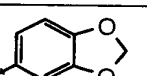
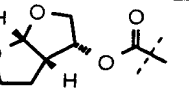
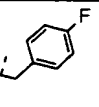

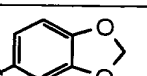
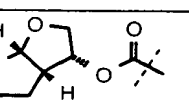
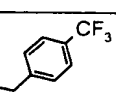

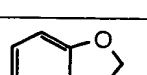
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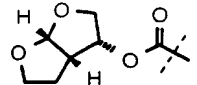
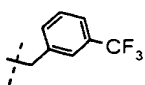

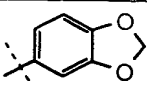
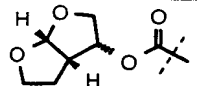
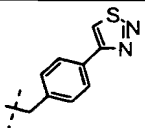

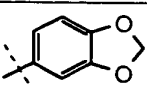
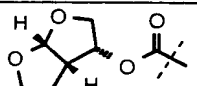
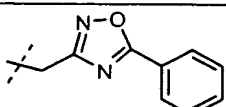

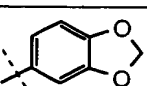
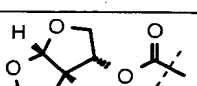
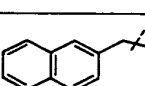

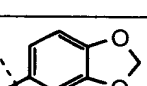
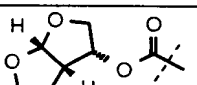
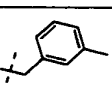

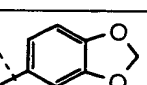
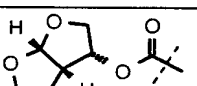
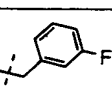

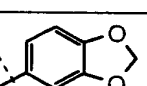
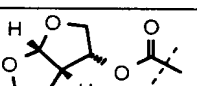
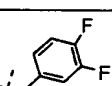

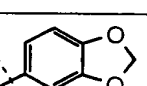
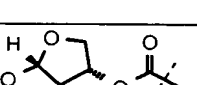
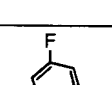

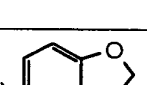
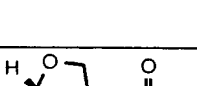
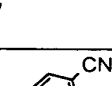

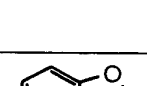
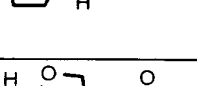


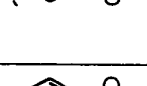
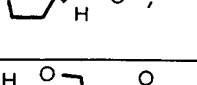
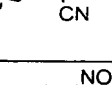

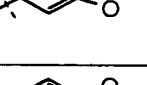
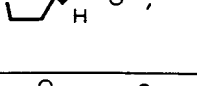
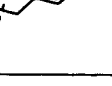

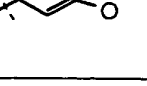
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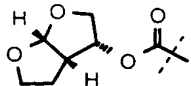

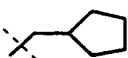
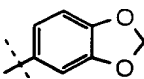
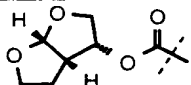

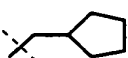
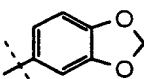
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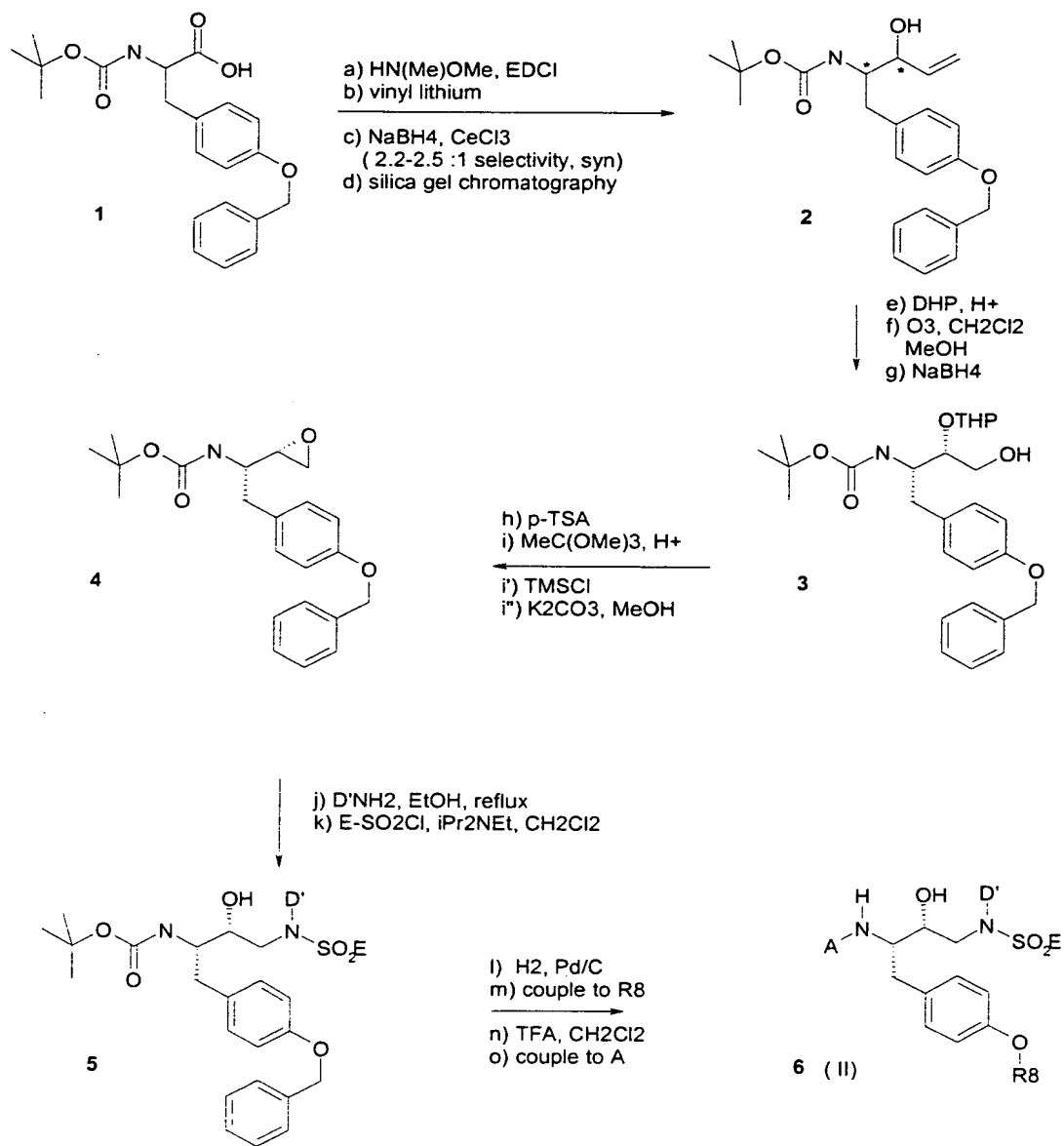
Preferred compound of the present invention are compound numbers: 18, 19, 20, 22, 24, 25, 26, 27, 31, 33, 35, 36, 38, 41, 43, 48, 49, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 68, 69, 71, 72, 73, 74, 202-204, 209, 213, 215, 217, 223, 227, 231, 233, 236, 237, 239, 243, 247, 250, 260, 263, 271, 281, 289, 293, 295, 304, 309, 317, 319, 320, 322, 334, 335, 348, 364, 367, 368, 375, 382, 383 and 396.

More preferred are compound numbers: 26, 27, 31, 33, 35, 36, 38, 41, 43, 48, 49, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 69, 71, 72, 73, 74, 209, 215, 227, 233, 237, 281, 289, 295, 304, 309, 322, 335, 364, 368, 382 and 383.

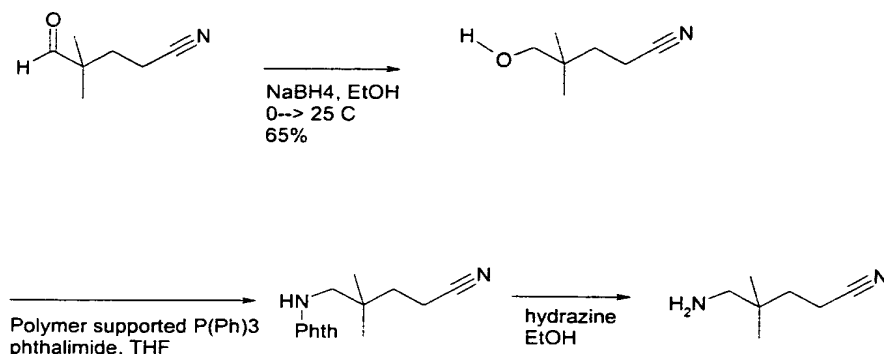
Most preferred are compound numbers: 54, 209, 237, 281, 295, 309, 367 and 368.

The compounds of the present invention can be readily prepared by techniques known in the art. Scheme I illustrates a general synthetic route to the compounds of this invention.

SCHEME I



SCHEME II



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In Step 1 of Scheme I, the bis protected amino acid is homologated through initial conversion to the Wienreb Amide (a) followed by alkylation with vinyl lithium (b) and stereoselective reduction (c). The diastereomers can be separated by silica gel chromatography (d). In step 2, the secondary alcohol is protected as a THP ether (e), as was found necessary for the oxidation step. The olefin was then oxidized to the aldehyde by ozone and the resulting ozonide reduced to the alcohol by sodium borohydride (steps f and g). After removal of the THP group (h) under acidic conditions the diol was converted to the epoxide (i, i' and i'') in one pot according to the method of Sharpless [K. B. Sharpless Tetrahedron 1992, 48 (35), pp. 10515-10530. The epoxide, 4, was then opened by H₂N-D' and further acylated in the presence of i-Pr₂Net by E-SO₂Cl to generate compounds of the formula schematically represented as 5. Alternate D' groups may be introduced at this point too. The synthesis of the D' as shown in compounds illustrated in Table II is shown in Scheme II.

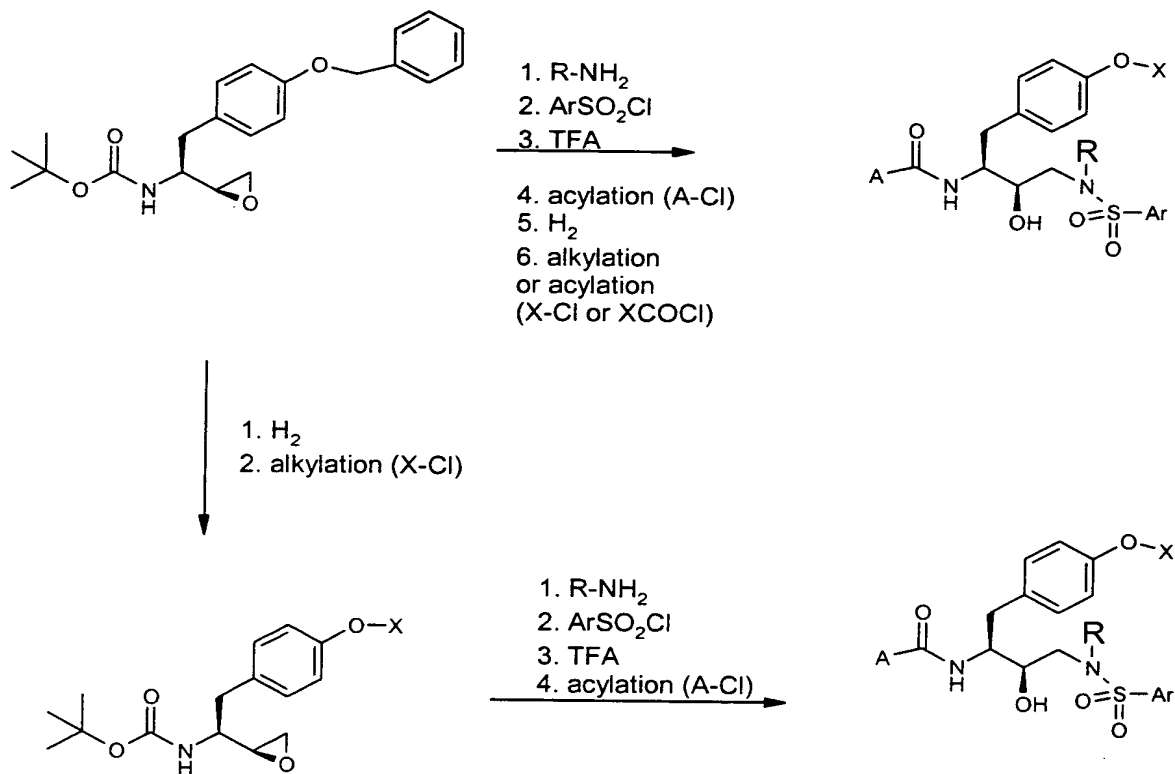
These compounds could then be further manipulated by removal of the Bn group and introduction

of a variety of R^8 groups by reacting with the
cooresponding alkyl halides. Further elaboration was
possible by removal of the t-Butyl carbonate (1) and
reintroduction of another group or carbamate designated
5 as A, to provide compounds represented as 6 (formula II).
We found that coupling as in reacton "m" was efficient
under the following general conditions: alkyl halide (R^8 -
Cl, 2.5 EQ. $CsCO_3$, dioxane, $80^\circ C$, 2-4 hours. Similar
alkylating conditions are reported in J. Med. Chem 1992,
10 1688 along with representative routes for the synthesis
of some R^8 -Cl intermediates. The coupling as illustrated
for bringing in A, step "o", was generally efficient
under the following conditions: activated p- NO_2 -phenyl
carbonate (p- NO_2 -O-A), i- Pr_2NEt , CH_2Cl_2 , RT, 12 hours. Use
15 of the activated succinate provides an alternative
coupling reagent (Succinate-A).

Alternatively, compounds of the present invention
can also be prepared according to Scheme III below.

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SCHEME III



Thus, the synthetic approach illustrated in Scheme I and Scheme III can be readily extended to produce other compounds of the present invention. The above synthetic schemes are not intended to comprise a comprehensive list of all means by which compounds described and claimed in this application may be synthesized. Further methods will be evident to those of ordinary skill in the art.

As discussed above, the novel compounds of the present invention are excellent ligands for aspartyl proteases, particularly HIV-1 and HIV-2 proteases. Accordingly, these compounds are capable of targeting and inhibiting late stage events in HIV replication, i.e., the processing of the viral polyproteins by HIV encoded proteases. Such compounds inhibit the proteolytic processing of viral polyprotein precursors by inhibiting

aspartyl protease. Because aspartyl protease is essential for the production of mature virions, inhibition of that processing effectively blocks the spread of virus by inhibiting the production of
5 infectious virions, particularly from chronically infected cells. Compounds according to this invention advantageously inhibit the ability of the HIV-1 virus to infect immortalized human T cells over a period of days, as determined by an assay of extracellular p24 antigen --
10 a specific marker of viral replication. Other anti-viral assays have confirmed the potency of these compounds.

The compounds of this invention may be employed in a conventional manner for the treatment of viruses, such as HIV and HTLV, which depend on aspartyl proteases for
15 obligatory events in their life cycle. Such methods of treatment, their dosage levels and requirements may be selected by those of ordinary skill in the art from available methods and techniques. For example, a compound of this invention may be combined with a
20 pharmaceutically acceptable adjuvant for administration to a virally-infected patient in a pharmaceutically acceptable manner and in an amount effective to lessen the severity of the viral infection.

Alternatively, the compounds of this invention may
25 be used in vaccines and methods for protecting individuals against viral infection over an extended period of time. The compounds may be employed in such vaccines either alone or together with other compounds of this invention in a manner consistent with the
30 conventional utilization of protease inhibitors in vaccines. For example, a compound of this invention may be combined with pharmaceutically acceptable adjuvants conventionally employed in vaccines and administered in

prophylactically effective amounts to protect individuals over an extended period time against HIV infection. As such, the novel protease inhibitors of this invention can be administered as agents for treating or preventing HIV
5 infection in a mammal.

The compounds of formula I, especially those having a molecular weight of less than about 700 g/mole, may be readily absorbed by the bloodstream of mammals upon oral administration. Compounds of formula I having a
10 molecular weight of less than about 600 g/mole are most likely to demonstrate oral availability. This surprisingly impressive oral availability makes such compounds excellent agents for orally-administered treatment and prevention regimens against HIV infection.

15 The compounds of this invention may be administered to a healthy or HIV-infected patient either as a single agent or in combination with other anti-viral agents which interfere with the replication cycle of HIV. By administering the compounds of this invention with other
20 anti-viral agents which target different events in the viral life cycle, the therapeutic effect of these compounds is potentiated. For instance, the co-administered anti-viral agent can be one which targets early events in the life cycle of the virus, such as cell
25 entry, reverse transcription and viral DNA integration into cellular DNA. Anti-HIV agents targeting such early life cycle events include, didanosine (ddI), alcitabine (ddC), d4T, zidovudine (AZT), polysulfated polysaccharides, sT4 (soluble CD4), ganciclovir,
30 dideoxycytidine, trisodium phosphonoformate, eflornithine, ribavirin, acyclovir, alpha interferon and trimenotrexate. Additionally, non-nucleoside inhibitors of reverse transcriptase, such as TIBO or nevirapine, may be

Combination therapies according to this invention exert a synergistic effect in inhibiting HIV replication because each component agent of the combination acts on a different site of HIV replication. The use of such combinations also advantageously reduces the dosage of a given conventional anti-retroviral agent which would be required for a desired therapeutic or prophylactic effect as compared to when that agent is administered as a monotherapy. These combinations may reduce or eliminate the side effects of conventional single anti-retroviral agent therapies while not interfering with the anti-retroviral activity of those agents. These combinations reduce potential of resistance to single agent therapies, while minimizing any associated toxicity. These combinations may also increase the efficacy of the conventional agent without increasing the associated toxicity. In particular, we have discovered that these compounds act synergistically in preventing the replication of HIV in human T cells. Preferred combination therapies include the administration of a compound of this invention with AZT, ddI, ddC or d4T.

Alternatively, the compounds of this invention may also be co-administered with other HIV protease inhibitors such as Ro 31-8959 (Roche), L-735,524 (Merck), XM 323 (Du-Pont Merck) and A-80,987 (Abbott) to increase the effect of therapy or prophylaxis against various viral mutants or members of other HIV quasi species.

We prefer administering the compounds of this invention as single agents or in combination with

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limited to, HTLV-I and HTLV-II. In addition, the compounds of this invention may also be used to inhibit other aspartyl proteases, and in particular, other human aspartyl proteases, including renin and aspartyl

5 proteases that process endothelin precursors.

Pharmaceutical compositions of this invention comprise any of the compounds of the present invention, and pharmaceutically acceptable salts thereof, with any pharmaceutically acceptable carrier, adjuvant or vehicle.

10 Pharmaceutically acceptable carriers, adjuvants and vehicles that may be used in the pharmaceutical compositions of this invention include, but are not limited to, ion exchangers, alumina, aluminum stearate, lecithin, serum proteins, such as human serum albumin,
15 buffer substances such as phosphates, glycine, sorbic acid, potassium sorbate, partial glyceride mixtures of saturated vegetable fatty acids, water, salts or electrolytes, such as protamine sulfate, disodium hydrogen phosphate, potassium hydrogen phosphate, sodium
20 chloride, zinc salts, colloidal silica, magnesium trisilicate, polyvinyl pyrrolidone, cellulose-based substances, polyethylene glycol, sodium carboxymethylcellulose, polyacrylates, waxes, polyethylene-polyoxypropylene-block polymers,
25 polyethylene glycol and wool fat.

The pharmaceutical compositions of this invention may be administered orally, parenterally, by inhalation spray, topically, rectally, nasally, buccally, vaginally or via an implanted reservoir. We prefer oral
30 administration or administration by injection. The pharmaceutical compositions of this invention may contain any conventional non-toxic pharmaceutically-acceptable carriers, adjuvants or vehicles. The term parenteral as

The pharmaceutical compositions may be in the form of a sterile injectable preparation, for example, as a sterile injectable aqueous or oleaginous suspension. This suspension may be formulated according to techniques known in the art using suitable dispersing or wetting agents (such as, for example, Tween 80) and suspending agents. The sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example, as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents that may be employed are mannitol, water, Ringer's solution and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose, any bland fixed oil may be employed including synthetic mono- or diglycerides. Fatty acids, such as oleic acid and its glyceride derivatives are useful in the preparation of injectables, as are natural pharmaceutically-acceptable oils, such as olive oil or castor oil, especially in their polyoxyethylated versions. These oil solutions or suspensions may also contain a long-chain alcohol diluent or dispersant such as Ph. Helv or a similar alcohol.

The pharmaceutical compositions of this invention may be orally administered in any orally acceptable dosage form including, but not limited to, capsules, tablets, and aqueous suspensions and solutions. In the case of tablets for oral use, carriers which are commonly used include lactose and corn starch. Lubricating

agents, such as magnesium stearate, are also typically added. For oral administration in a capsule form, useful diluents include lactose and dried corn starch. When aqueous suspensions are administered orally, the active
5 ingredient is combined with emulsifying and suspending agents. If desired, certain sweetening and/or flavoring and/or coloring agents may be added.

The pharmaceutical compositions of this invention may also be administered in the form of suppositories for
10 rectal administration. These compositions can be prepared by mixing a compound of this invention with a suitable non-irritating excipient which is solid at room temperature but liquid at the rectal temperature and therefore will melt in the rectum to release the active
15 components. Such materials include, but are not limited to, cocoa butter, beeswax and polyethylene glycols.

Topical administration of the pharmaceutical compositions of this invention is especially useful when the desired treatment involves areas or organs readily
20 accessible by topical application. For application topically to the skin, the pharmaceutical composition should be formulated with a suitable ointment containing the active components suspended or dissolved in a carrier. Carriers for topical administration of the
25 compounds of this invention include, but are not limited to, mineral oil, liquid petroleum, white petroleum, propylene glycol, polyoxyethylene polyoxypropylene compound, emulsifying wax and water. Alternatively, the pharmaceutical composition can be formulated with a
30 suitable lotion or cream containing the active compound suspended or dissolved in a carrier. Suitable carriers include, but are not limited to, mineral oil, sorbitan monostearate, polysorbate 60, cetyl esters wax, cetearyl

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alcohol, 2-octyldodecanol, benzyl alcohol and water. The pharmaceutical compositions of this invention may also be topically applied to the lower intestinal tract by rectal suppository formulation or in a suitable enema
5 formulation. Topically-transdermal patches are also included in this invention.

The pharmaceutical compositions of this invention may be administered by nasal aerosol or inhalation. Such compositions are prepared according to techniques well-
10 known in the art of pharmaceutical formulation and may be prepared as solutions in saline, employing benzyl alcohol or other suitable preservatives, absorption promoters to enhance bioavailability, fluorocarbons, and/or other solubilizing or dispersing agents known in the art.

15 Dosage levels of between about 0.01 and about 100 mg/kg body weight per day, preferably between about 0.5 and about 50 mg/kg body weight per day of the active ingredient compound are useful in the prevention and treatment of viral infection, including HIV infection.
20 Typically, the pharmaceutical compositions of this invention will be administered from about 1 to about 5 times per day or alternatively, as a continuous infusion. Such administration can be used as a chronic or acute therapy. The amount of active ingredient that may be
25 combined with the carrier materials to produce a single dosage form will vary depending upon the host treated and the particular mode of administration. A typical preparation will contain from about 5% to about 95% active compound (w/w). Preferably, such preparations
30 contain from about 20% to about 80% active compound.

Upon improvement of a patient's condition, a maintenance dose of a compound, composition or combination of this invention may be administered, if

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necessary. Subsequently, the dosage or frequency of
administration, or both, may be reduced, as a function of
the symptoms, to a level at which the improved condition
is retained when the symptoms have been alleviated to the
5 desired level, treatment should cease. Patients may,
however, require intermittent treatment on a long-term
basis upon any recurrence of disease symptoms.

As the skilled artisan will appreciate, lower or
higher doses than those recited above may be required.
10 Specific dosage and treatment regimens for any particular
patient will depend upon a variety of factors, including
the activity of the specific compound employed, the age,
body weight, general health status, sex, diet, time of
administration, rate of excretion, drug combination, the
15 severity and course of the infection, the patient's
disposition to the infection and the judgment of the
treating physician.

The compounds of this invention are also useful as
commercial reagents which effectively bind to aspartyl
20 proteases, particularly HIV aspartyl protease. As
commercial reagents, the compounds of this invention, and
their derivatives, may be used to block proteolysis of a
target peptide or may be derivatized to bind to a stable
resin as a tethered substrate for affinity chromatography
25 applications. These and other uses which characterize
commercial aspartyl protease inhibitors will be evident
to those of ordinary skill in the art.

As used herein, the compounds according to the
invention are defined to include pharmaceutically
30 acceptable derivatives or prodrugs thereof. A
"pharmaceutically acceptable derivative" or
"pharmaceutically acceptable prodrug" means any
pharmaceutically acceptable salt, ester, salt of an

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Salts derived from appropriate bases include alkali metal (e.g. sodium), alkaline earth metal (e.g.,

magnesium), ammonium and NW_4 (wherein W is C_{1-4} alkyl). Physiologically acceptable salts of a hydrogen atom or an amino group include salts or organic carboxylic acids such as acetic, lactic, tartaric, malic, isethionic, 5 lactobionic and succinic acids; organic sulfonic acids such as methanesulfonic, ethanesulfonic, benzenesulfonic and p-toluenesulfonic acids and inorganic acids such as hydrochloric, sulfuric, phosphoric and sulfamic acids. Physiologically acceptable salts of a compound with a 10 hydroxy group include the anion of said compound in combination with a suitable cation such as Na^+ , NH_4^+ , and NW_4^+ (wherein W is a C_{1-4} alkyl group).

Pharmaceutically acceptable salts include salts of organic carboxylic acids such as ascorbic, acetic, 15 citric, lactic, tartaric, malic, maleic, isothionic, lactobionic, p-aminobenzoic and succinic acids; organic sulphonic acids such as methanesulphonic, ethanesulphonic, benzenesulphonic and p-toluenesulphonic acids and inorganic acids such as hydrochloric, 20 sulphuric, phosphoric, sulphamic and pyrophosphoric acids.

For therapeutic use, salts of the compounds according to the invention will be pharmaceutically acceptable. However, salts of acids and bases which are 25 non-pharmaceutically acceptable may also find use, for example, in the preparation or purification of a pharmaceutically acceptable compound.

Preferred salts include salts formed from hydrochloric, sulfuric, acetic, succinic, citric and 30 ascorbic acids.

Preferred esters of the compounds according to the invention are independently selected from the following groups: (1) carboxylic acid esters obtained by

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esterification of the hydroxy groups, in which the non-carbonyl moiety of the carboxylic acid portion of the ester grouping is selected from straight or branched chain alkyl (for example, acetyl, n-propyl, t-butyl, or n-butyl), alkoxyalkyl (for example, methoxymethyl), aralkyl (for example, benzyl), aryloxyalkyl (for example, phenoxymethyl), aryl (for example, phenyl optionally substituted by, for example, halogen, C₁₋₄alkyl, or C₁₋₄alkoxy or amino); (2) sulfonate esters, such as alkyl- or aralkylsulfonyl (for example, methanesulfonyl); (3) amino acid esters (for example, L-valyl or L-isoleucyl); (4) phosphonate esters and (5) mono-, di- or triphosphate esters. The phosphate esters may be further esterified by, for example, a C₁₋₂₀ alcohol or reactive derivative thereof, or by a 2,3-di (C₆₋₂₄)acyl glycerol.

In such esters, unless otherwise specified, any alkyl moiety present advantageously contains from 1 to 18 carbon atoms, particularly from 1 to 6 carbon atoms, more particularly from 1 to 4 carbon atoms. Any cycloalkyl moiety present in such esters advantageously contains from 3 to 6 carbon atoms. Any aryl moiety present in such esters advantageously comprises a phenyl group.

Any reference to any of the above compounds also includes a reference to a pharmaceutically acceptable salts thereof.

The compounds according to the invention are especially useful for the treatment of AIDS and related clinical conditions such as AIDS related complex (ARC), progressive generalized lymphadenopathy (PGL), Kaposi's sarcoma, thrombocytopenic purpura, AIDS-related neurological conditions such as AIDS dementia complex, multiple sclerosis or tropical paraperesis, and also

anti-HIV antibody-positive and HIV-positive conditions, including such conditions in asymptomatic patients.

In a further aspect of the invention there are provided the compounds according to the invention for use
5 in medical therapy particularly for the treatment or prophylaxis of viral infections such as HIV infections.

According to another aspect, the present invention provides a method for the treatment or prevention of the symptoms or effects of a viral
10 infection in an infected animal, for example, a mammal including a human, which comprises treating said animal with a therapeutically effective amount of a compound according to the invention. According to a particular embodiment of this aspect of the invention, the viral
15 infection is an HIV infection. A further aspect of the invention includes a method for the treatment or prevention of the symptoms or effects of an HBV infection.

The compounds according to the invention may also be
20 used in adjuvant therapy in the treatment of HIV infections or HIV-associated symptoms or effects, for example Kaposi's sarcoma.

The present invention further provides a method for the treatment of a clinical condition in an animal,
25 for example, a mammal including a human which clinical condition includes those which have been discussed in the introduction hereinbefore, which comprises treating said animal with a therapeutically effective amount of a compound according to the invention. The present
30 invention also includes a method for the treatment or prophylaxis of any of the aforementioned infections or conditions.

Reference herein to treatment extends to prophylaxis as well as the treatment of established infections or symptoms.

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The above compounds according to the invention
5 and their pharmaceutically acceptable derivatives may be employed in combination with other therapeutic agents for the treatment of the above infections or conditions. Combination therapies according to the present invention comprise the administration of at least one compound of
10 the formula (I) or a pharmaceutically acceptable derivative thereof and at least one other pharmaceutically active ingredient. The active ingredient(s) and pharmaceutically active agents may be administered simultaneously in either the same or
15 different pharmaceutical formulations or sequentially in any order. The amounts of the active ingredient(s) and pharmaceutically active agent(s) and the relative timings of administration will be selected in order to achieve the desired combined therapeutic effect. Preferably the
20 combination therapy involves the administration of one compound according to the invention and one of the agents mentioned herein below.

Sub
25 Examples of such further therapeutic agents include agents that are effective for the treatment of viral infections or associated conditions such as (1 alpha, 2 beta, 3 alpha)-9-[2,3-bis(hydroxymethyl) cyclobutyl]guanine [(-)BHCg, SQ-34514], oxetanocin-G (3,4-bis-(hydroxymethyl)-2-oxetanosyl]guanine), acyclic nucleosides (e.g. acyclovir, valaciclovir, famciclovir,
30 ganciclovir, penciclovir), acyclic nucleoside phosphonates (e.g. (S)-1-(3-hydroxy-2-phosphonyl-methoxypropyl)cytosine (HPMPC) and PMEA analogs thereof, ribonucleotide reductase inhibitors such as 2-

acetylpyridine 5-[(2-chloroanilino)thiocarbonyl)
thiocarbonohydrazone, 3'-azido-3'-deoxythymidine, other
2',3'-dideoxynucleosides such as 2',3'-dideoxycytidine,
2',3'-dideoxyadenosine, 2',3'-dideoxyinosine, 2',3'-
5 didehydrothymidine, protease inhibitors such as
indinavir, ritonavir, nelfinavir, [3S-[3R*(1R*, 2S*)]]-
[3[[[(4-aminophenyl)sulfonyl](2-methylpropyl)amino]-2-
hydroxy-1-(phenylmethyl)propyl]-tetrahydro-3-furanyl
ester (141W94), oxathiolane nucleoside analogues such as
10 (-)-cis-1-(2-hydroxymethyl)-1,3-oxathiolane 5-yl)-
cytosine (lamivudine) or cis-1-(2-(hydroxymethyl)-1,3-
oxathiolan-5-yl)-5-fluorocytosine (FTC), 3'-deoxy-3'-
fluorothymidine, 5-chloro-2',3'-dideoxy-3'-fluorouridine,
(-)-cis-4-[2-amino-6-(cyclopropylamino)-9H-purin-9-yl]-2-
15 cyclopentene-1-methanol, ribavirin, 9-[4-hydroxy-2-
(hydroxymethyl)but-1-yl]-guanine (H2G), tat inhibitors
such as 7-chloro-5-(2-pyrrolyl)-3H-1,4-benzodiazepin-2-
(H)one (Ro5-3335), 7-chloro-1,3-dihydro-5-(1H-pyrrol-
2-yl)-3H-1,4-benzodiazepin-2-amine (Ro24-7429),
20 interferons such as α -interferon, renal excretion
inhibitors such as probenecid, nucleoside transport
inhibitors such as dipyrindamole; pentoxifylline, N-
acetylcysteine (NAC), Procysteine, α -trichosanthin,
phosphonoformic acid, as well as immunomodulators such as
25 interleukin II or thymosin, granulocyte macrophage colony
stimulating factors, erythropoietin, soluble CD₄ and
genetically engineered derivatives thereof, or non-
nucleoside reverse transcriptase inhibitors (NNRTIs) such
as nevirapine (BI-RG-587), zalcitabine (α -APA) and
30 delavuridine (BHAP), and phosphonoformic acid, and 1,4-
dihydro-2H-3,1-benzoxazin-2-ones NNRTIs such as (-)-6-
chloro-4-cyclopropylethynyl-4-trifluoromethyl-1,4-
dihydro-2H-3,1-benzoxazin-2-one (L-743,726 or DMP-266),

and quinoxaline NNRTIs such as isopropyl (2S)-7-fluoro-3,4-dihydro-2-ethyl-3-oxo-1(2H)-quinoxalinecarboxylate (HBY1293).

More preferably the combination therapy
5 involves the administration of one of the above mentioned agents and a compound within one of the preferred or particularly preferred sub-groups within formula (I) as described above. Most preferably the combination therapy involves the joint use of one of the above named agents
10 together with one of the compounds of formula (I) specifically named herein.

The present invention further includes the use of a compound according to the invention in the manufacture of a medicament for simultaneous or sequential
15 administration with at least one other therapeutic agent, such as those defined herein before.

In order that this invention may be more fully understood, the following examples are set forth. These examples are for the purpose of illustration only and are
20 not to be construed as limiting the scope of the invention in any way. Compounds for which experimentals are not shown may be made through similar methodologies.

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Example 1

**Synthesis of BOC Benzyl Tyrosine Based
Weinreb Amide (Scheme 1, Step a)**

5 N-t-BOC-O-Benzyl-L-Tyrosine(1, Sigma) (25 g, 67.3
mmol) was combined with anhydrous DMF (200 ml) and cooled
to 0°C under a N₂ atmosphere. HOBt (15.5 g, 114.4 mmol,
1.7 eq.) and EDC (15.5 g, 80.8 mmol, 1.2 eq.) were added
as solids and stirred to dissolve. Diisopropylethylamine
10 (17.6 ml, 101 mmol, 1.5 eq.) and 4-dimethylaminopyridine
(0.001 g) were added and the reaction was stirred for 50
minutes at 0°C. N,O-Dimethylhydroxylamine
hydrochloride(8.5 g, 87.5 mmol, 1.3 eq.) was added as a
solid and the reaction was stirred for 10 minutes at 0°C
15 then allowed to warm to room temperature and stirred
overnight. After 18 hours at room temperature, the
reaction was cooled to 0°C, and quenched with 200 mL of a
5% sodium bicarbonate solution. The reaction was
extracted twice with EtOAc. The combined organics were
20 washed with five times with water, and then brine, dried
over MgSO₄, filtered and the solvent was removed *in vacuo*.
The yield was 28g of amide which was used as is.
HPLC (5-100% CH₃CN/water) showed 1 peak at 10.77 min and
the NMR (CDCl₃) was consistent with the expected
25 structure.

Example 2

**BOC Benzyl Tyrosine derived Vinyl Ketone (Scheme 1, Step
b)**

30 N-t-BOC-O-Benzyl-L-Tyrosine Weinreb amide (18.8 g,
45.3 mmol) was combined with anhydrous THF (200 ml) and
cooled to -78°C. A vinyl lithium solution (2.3 M, 50 ml,

2.5 eq.) was added via addition funnel dropwise over 20 minutes at -78°C . Ten mL of anhydrous THF was added to rinse the funnel. The reaction was stirred at -78°C under a N_2 atmosphere. HPLC at 1.5 hours showed the reaction was ~50% complete. Another 1.0 eq (20ml) of vinyl lithium was added over 10 minutes at -78°C and washed in with 15mL of THF. The reaction stirred overnight at -78°C . After 18 hours, HPLC showed ~15% Weinreb amide left. Another 0.2 eq (4 ml) of vinyl lithium was added at -78°C . After 26 hours at -78°C , the reaction was quenched with a slow addition of 300mL of 1N HCl. The reaction was partitioned between EtOAc and water. The aqueous phase was extracted with EtOAc. The combined organics were washed consecutively with saturated bicarbonate solution and brine, dried over MgSO_4 , filtered and the solvent was removed in vacuo. The yield was 19.9 g crude material.

The material was purified by flash chromatography (gradient: CH_2Cl_2 to 10%EtOAc/ CH_2Cl_2) to give 13.5g (78%) of pure material. HPLC (5-100% CH_3CN /water) showed 1 peak at 13.37 min and the LC/MS showed 1 peak with an $\text{M}+\text{H} = 382.4$ for the desired compound.

Example 3

BOC Benzyl Tyrosine derived Allyl Alcohol (Scheme 1, step c, compound 2)

N-t-BOC-O-benzyl-L-tyrosine vinyl ketone (13.5 g, 35.4 mmol) was combined with methanol (120 ml) and Methylene Chloride (30 mL) and cooled to 0°C . Cerium chloride heptahydrate (14.5 g, 39 mmol, 1.1 eq.) was then added as a solid. The reaction was stirred at 0°C

for 5 minutes and then cooled to -78°C . A solution of sodium borohydride (2.0 g, 53.1 mmol, 1.5 eq) in 40 mL of MeOH was cooled to -78°C and cannulated into the reaction dropwise over 40 minutes. The reaction became a thick
5 white suspension and 50 mL of MeOH was added to aid stirring. The reaction was stirred at -78°C for 1.5 hours and was then quenched at -78°C with 150 mL of a saturated ammonium chloride solution. The reaction was extracted three times with EtOAc, and the combined
10 organics were washed with saturated bicarbonate solution, followed by brine, dried over Na_2SO_4 , filtered and the solvent was removed in vacuo to give 13.6 g crude material.

Proton NMR (CDCl_3) shows a 7:3 ratio of
15 diastereomers. The material was purified via chromatography (4:5:1, Hexanes: CH_2Cl_2 :EtOAc) to give 5.4 g of desired material (83:17 ratio of diastereomers) as well as 7.3 g of allyl alcohol as a mix of diastereomers to be repurified. Proton NMR (CDCl_3) was consistent with
20 structure for the desired material.

Example 4

BOC Benzyl Tyrosine Allyl Alcohol

(THP protected) Scheme 1, step e:

25

BOC benzyl tyrosine allyl alcohol (1.59 g, 4.1 mmole) was dissolved in 10 mL of anhydrous CH_2Cl_2 . Dihydropyran (500 μL , 5.4 mmol, 1.3eq.) and pyridinium paratoluenesulfonate (210 mg, 0.8 mmol, 0.2 eq.) was
30 added and the reaction was stirred at room temperature under a N_2 atmosphere. After 19 hours, the solvent was removed in vacuo and the residue was partitioned between

EtOAc and a 10% citric acid solution. The organics were separated, washed with brine, and then saturated bicarbonate solution, dried over Na_2SO_4 , filtered and the solvent was removed *in vacuo* to give 1.98 g crude material as a white solid.

The material was purified via chromatography (25% EtOAc/Hexanes, (with 0.5ml NEt_3/L)) to give 1.85 g (96%) of desired material. NMR (CDCl_3) was consistent with structure as a mix of diastereomers (1:1) at THP alcohol.

Example 5

BOC Benzyl Tyrosine Diol (THP protected)
(Scheme 1, steps f and g, compound 3)

THP protected BOC benzyl tyrosine allyl alcohol (1.5 g, 3.2 mmol) was dissolved in methanol (5 ml) and methylene chloride (20 mL) and cooled to -78°C . Ozone was bubbled into the stirred solution for 1.5 hours at -78°C . The solution was then flushed with nitrogen to remove the ozone. Sodium borohydride (920 mg, 25.6 mmol, 8 eq.) was then added in small portions over 5 minutes at -78°C . Methanol (35 mL) was added and the reaction was stirred at 78°C for 5 minutes; then was slowly warmed to 0°C . Vigorous bubbling began at $\sim -20^\circ\text{C}$. After 1.5 hours at 0°C the reaction was quenched with saturated bicarbonate solution and extracted with CH_2Cl_2 . The combined organics were washed with brine, dried over Na_2SO_4 , filtered and the solvent was removed *in vacuo* to give 1.49 g crude material.

The material was purified via chromatography (EtOAc/Hexanes, 20%-30%-40%, containing 0.5ml NEt_3/L) to give 1.15 g (76%) of desired material. HPLC (5-100%

CH₃CN/water) showed 1 peak at 13.81 min and the proton NMR (CDCl₃) was consistent with structure as a mix of diastereomers (~1:1) at the THP alcohol.

5

Example 6

Epoxide (4)

THP-protected diol (3) (0.40 g, 0.85 mmol) was combined with a catalytic amount of *p*-toluenesulfonic acid (0.004 g) in methanol (20 mL) under a N₂ atmosphere. After stirring at room temperature for ca. 15 minutes, the initial white suspension dissolved completely. Stirring at room temperature was continued for ca. 1 hour, after which time complete disappearance of starting material (3) was confirmed by TLC. The solvent was removed *in vacuo* to give diol as a white solid, combined with residual *p*-toluenesulfonic acid. The crude diol was dissolved in anhydrous dichloromethane (15 mL), and trimethylorthoacetate (0.130 mL, 1.02 mmol) was added dropwise with stirring. Upon addition of trimethylorthoacetate, the cloudy solution became colourless. After stirring at room temperature for ca. 1 hour, TLC again indicated complete disappearance of starting material. The solvent was removed *in vacuo* to give the desired cyclic orthoacetate as a viscous white oil, which was re-dissolved in anhydrous DCM (15 mL). Trimethylsilyl chloride (0.129 mL, 1.02 mmol) was added dropwise with stirring. After 1.5 hours at room temperature generally no further starting material remained. The solvent was removed *in vacuo* to give the desired chloroacetates as a yellow oil, which was dissolved in methanol (20 mL). Cesium carbonate (0.48 g, 1.48 mmol) was added in one portion, and the solution was

stirred at room temperature for 2 hours, after which time, TLC confirmed the absence of any starting material. The solvent was removed *in vacuo* to give a pale yellow oil, which was partitioned between saturated aqueous ammonium chloride solution (30 mL) and DCM (30 mL). The organic layer was taken, and the aqueous layer re-extracted with DCM (2 x 30 mL). The combined organic extracts were dried over MgSO_4 , and the solvent removed *in vacuo* to give crude epoxide (4) as a yellow oil. This material was either used in the crude form, or could be purified by flash column chromatography (3:7 ethyl acetate/hexane to 4:1 ethyl acetate/methanol) to give epoxide (4) as a white solid: $R_f = 0.60$ (3:7 ethyl acetate/hexane); $^1\text{H NMR}$ (CDCl_3) 7.49-7.29 (5H, m), 7.14 (2H, d, $J = 8.3$ Hz), 6.93 (2H, d, $J = 8.3$ Hz), 5.05 (2H, s), 4.44 (1H, br. s), 3.64 (1H, br. s), 2.97-2.86 (2H, m), 2.85-2.72 (3H, m), 1.39 (9H, s); coupled LCMS showed the product as a single major peak with m/z 370 $[\text{M}+\text{H}]^+$ or m/z 312 $[\text{M}+\text{H}-\text{tBu}]^+$ at R_T 2.84 min; HPLC (205 nm) over an extended 20 minute run time showed the crude material to be a 9:1 mixture of diastereoisomeric epoxides at R_T s of 14.2 min. (*major diastereoisomer*) & 14.3 min. (*minor diastereoisomer*).

25

Example 7

Isobutylamino Alcohol (Scheme 1, step j)

Crude epoxide (4) (0.37 g, 0.85 mmol) was combined with isobutylamine (*large excess*, 4 mL) in ethanol (4 mL) under a N_2 atmosphere. The reaction mixture was heated to reflux with stirring for 2.5 hours. The solvent was removed *in vacuo* to give a pale yellow oily residue. Trituration with hexane gave the isobutylamino alcohol

(0.285 g, 75%) as a white solid: R_f = 0.05 (3:7 ethyl acetate/hexane); ^1H NMR (CDCl_3) 7.47-7.29 (5H, m), 7.15 (2H, d, J = 8.5 Hz), 6.91 (2H, d, J = 8.5 Hz), 5.04 (2H, s), 4.69 (1H, br. d, J = 8.8 Hz), 3.76 (1H, br. s), 3.49-3.40 (1H, m), 2.91 (1H, dd, J = 14.1, 4.7 Hz), 2.87-2.77 (1H, m), 2.67 (2H, d, J = 4.7 Hz), 2.40 (2H, d, J = 6.7 Hz), 1.71 (1H, septet, J = 6.7 Hz), 1.36 (9H, s), 0.91 (3H, d, J = 6.7 Hz), 0.90 (3H, d, J = 6.7 Hz), OH and NH signals not observed; distinct *minor diastereoisomer* signals observed: 4.55 (1H, d, J = 8.7 Hz), 2.49 (2H, d, J = 6.6 Hz); NMR integration showed the triturated product to be a 9:1 mixture of diastereoisomeric isobutylamino alcohols; coupled LCMS showed the product as a single major peak with m/z 443 $[\text{M}+\text{H}]^+$ at R_T 2.41 min.

Example 8

Boc methylenedioxybenzenesulfonamide benzyl ether (Scheme 1, Compound 5e)

Isobutylamino alcohol (0.170 g, 0.38 mmol) was combined with methylenedioxybenzenesulfonyl chloride (0.085 g, 0.38 mmol) in anhydrous DCM (5 mL) under a N_2 atmosphere. The solution was cooled to 0 °C using an ice bath, and diisopropylethylamine (0.20 mL, 1.20 mmol) was added dropwise, and the reaction was allowed to warm to room temperature with stirring for 4 hours. The solvent was removed *in vacuo*, and the resultant pale yellow oil was purified by flash column chromatography (3:7 ethyl acetate/hexane) to give Boc methylenedioxybenzenesulfonamide benzyl ether (0.185 g, 75%) as a white foam: R_f = 0.30 (3:7 ethyl acetate/hexane); ^1H NMR (CDCl_3) 7.47-7.29 (6H, m), 7.21-7.11 (3H, m), 6.92 (2H, d, J = 8.4 Hz), 6.88 (1H, d, J =

8.2 Hz), 6.07 (2H, s), 5.04 (2H, s), 4.67-4.58 (1H, m), 3.97-3.85 (1H, m), 3.82-3.56 (2H, m), 3.10-3.02 (2H, m), 2.98-2.72 (4H, m), 1.84 (1H, septet, $J = 6.5$ Hz), 1.36 (9H, s), 0.91 (3H, d, $J = 6.5$ Hz), 0.87 (3H, d, $J = 6.5$ Hz); coupled LCMS showed the product as a single major peak with m/z 627 $[M+H]^+$ at R_T 3.16 min.

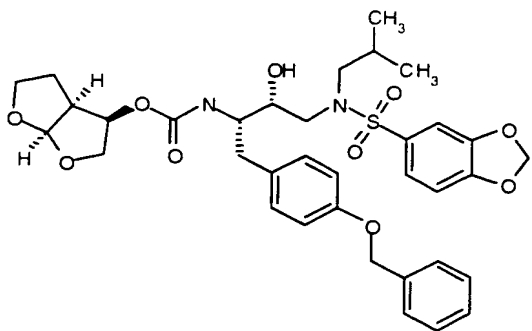
Example 9

Boc *m*-nitrobenzenesulfonamide

10 Benzyl Ether (Scheme 1, Compound 5c)

The isobutylamino alcohol (0.059 g, 0.13 mmol), as made in Scheme I by the addition of *i*-BuNH₂ to compound 4, was combined with *m*-nitrobenzenesulfonyl chloride (0.044 g, 0.20 mmol) in anhydrous DCM (2 mL) under a N₂ atmosphere. Diisopropylethylamine (0.070 mL, 0.40 mmol) was added dropwise, and the reaction was stirred at room temperature for 48 hours. The solvent was removed in vacuo, and the resultant yellow oil was purified by flash column chromatography (3:7 ethyl acetate/hexane) to give Boc *m*-nitrobenzenesulfonamide benzyl (5) (0.050 g, 61%) as a colourless oil: $R_f = 0.31$ (3:7 ethyl acetate/hexane); ¹H NMR (CDCl₃) 8.63 (1H, d, $J = 1.8$ Hz), 8.41 (1H, d, $J = 8.1$ Hz), 8.10 (1H, d, $J = 6.8$ Hz), 7.72 (1H, t, $J = 7.9$ Hz), 7.50-7.28 (5H, m), 7.15 (2H, d, $J = 8.6$ Hz), 6.92 (2H, d, $J = 8.6$ Hz), 5.00 (2H, s), 4.65-4.55 (1H, m), 3.99-3.84 (1H, m), 3.83-3.74 (1H, m), 3.74-3.64 (1H, m), 3.21 (2H, d, $J = 5.4$ Hz), 3.01 (2H, d, $J = 7.2$ Hz), 2.93-2.80 (2H, m), 1.88 (1H, septet, $J = 6.4$ Hz), 1.36 (9H, s), 0.91-0.75 (6H, m).

Example 10

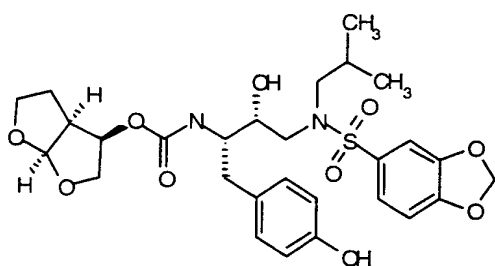


**Bis-THF methylenedioxybenzenesulfonamide benzyl ether
(46)**

- 5 Boc methylenedioxybenzenesulfonamide benzyl ether (0.040 g, 0.064 mmol) was dissolved in DCM (2 mL). Trifluoroacetic acid (1 mL) was added dropwise, and the reaction was stirred at room temperature for 1 hour. The solvent was removed *in vacuo*, and the resultant orange
- 10 oil was dissolved in DCM (1.5 mL) and the solution cooled to 0 °C using an ice bath. Diisopropylethylamine (0.33 mL, 1.92 mmol) was added dropwise with stirring, followed by (3R, 3aS, 6aR) hexahydrofuro[2,3-*b*]furan-2-yl 4-nitrophenyl carbonate (0.021 g, 0.071 mmol) in one
- 15 portion as a solid. After 5 minutes, the ice bath was removed and the reaction mixture was allowed to warm to room temperature with stirring overnight. The solvent was removed *in vacuo*, and the resultant yellow oil was purified by flash column chromatography (1:1 ethyl
- 20 acetate/hexane) to give *bis*-THF methylenedioxybenzenesulfonamide benzyl ether (46) (0.035 g, 80%) as a white foam: R_f = 0.31 (1:1 ethyl acetate/hexane); ^1H NMR (CDCl_3) 7.45-7.29 (6H, m), 7.17 (1H, d, J = 1.8 Hz), 7.13 (2H, d, J = 8.3 Hz), 6.90 (2H, d, J = 8.3 Hz), 6.94-6.86 (1H, m), 6.07 (2H, s), 5.66 (1H, d, J = 5.2 Hz), 5.10-4.98 (1H, m), 5.02 (2H, s), 4.94 (1H, d, J = 8.5 Hz), 3.96 (1H, dd, J = 9.6, 6.4 Hz),

3.89-3.78 (3H, m), 3.76-3.65 (2H, m), 3.18-3.09 (1H, m),
3.04-2.85 (4H, m), 2.82-2.70 (2H, m), 1.90-1.77 (1H, m),
1.73-1.48 (2H, m), 0.93 (3H, d, J = 6.5 Hz), 0.89 (3H, d,
J = 6.5 Hz), OH signal not observed; coupled LCMS showed
5 the product as a single major peak with m/z 683 $[M+H]^+$;
HPLC (205 nm) showed the material as a single major peak
at R_T of 2.73 min. (purity = 96%).

Example 11



10

***Bis*-THF methylenedioxybenzenesulfonamide free phenol (47)**

A solution of *bis*-THF methylenedioxybenzenesulfonamide
benzyl ether (46) (0.022 g, 0.032 mmol) and 10% palladium
15 on carbon (wet; Degussa variant) (0.008 g) in degassed
ethyl acetate (10 mL) was stirred under a balloon of
hydrogen. The reaction was monitored by TLC, and after 20
hours the starting material (47) had not been consumed. A
fresh portion of 10% palladium on carbon (wet; Degussa
20 variant) (0.008 g) was added to the reaction mixture, and
the reaction stirred under a balloon of hydrogen for a
further 4.5 hours. The reaction mixture was filtered
through a pad of celite, and the filtrate was dried *in*
vacuo to give a pale yellow oil. Flash column
25 chromatography (1:1 ethyl acetate/hexane) gave *bis*-THF
methylenedioxybenzenesulfonamide free phenol (47) (0.008
g, 42%) as a colourless oil: R_f = 0.44 (1:1 ethyl
acetate/hexane); 1H NMR ($CDCl_3$); 7.33 (1H, dd, J = 8.2,

1.7 Hz), 7.17 (1H, s), 7.07 (2H, d, J = 8.1 Hz), 6.90 (1H, d, J = 8.2 Hz), 6.74 (2H, d, J = 8.1 Hz), 6.09 (2H, s), 5.66 (1H, d, J = 5.2 Hz), 5.10-5.00 (2H, m), 4.05-3.68 (7H, m), 3.18-3.07 (1H, m), 3.06-2.90 (4H, m), 2.87-2.68 (2H, m), 1.89-1.48 (3H, m), 0.93 (3H, d, J = 6.6 Hz), 0.89 (3H, d, J = 6.6 Hz); OH signal not observed; coupled LCMS showed the product as a single major peak with m/z 593 $[M+H]^+$; HPLC (205 nm) showed the material as a single major peak at R_T of 1.94 min. (purity = 98%).

Example 12

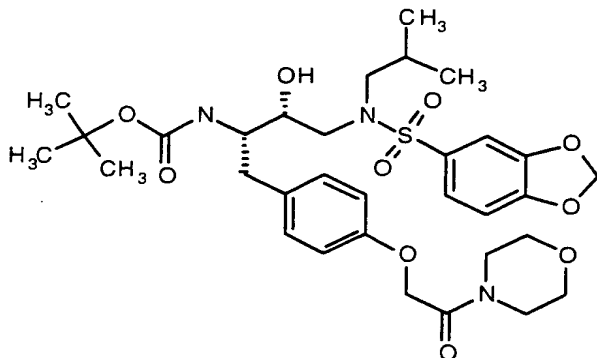
Boc methylenedioxybenzenesulfonamide free phenol (Scheme 1, Compound 6e)

A solution of Boc methylenedioxybenzenesulfonamide benzyl ether (0.063 g, 0.101 mmol) and 10% palladium on carbon (wet; Degussa variant) (0.024 g) in degassed ethyl acetate (10 mL) was stirred under a balloon of hydrogen. The reaction was monitored by TLC, and after 2 hours the starting material had not been consumed. A fresh portion of 10% palladium on carbon (wet; Degussa variant) (0.024 g) was added to the reaction mixture, and the reaction stirred under a balloon of hydrogen for a further 5 hours. The reaction mixture was filtered through a pad of celite, and the filtrate was dried *in vacuo* to give a colourless oil. Flash column chromatography (1:1 ethyl acetate/hexane) gave Boc methylenedioxybenzenesulfonamide free phenol (6) (0.036 g, 60%) as a white foam: R_f = 0.74 (1:1 ethyl acetate/hexane); 1H NMR ($CDCl_3$); 7.32 (1H, d, J = 8.3 Hz), 7.17 (1H, s), 7.12-7.02 (2H, m), 6.88 (1H, d, J = 8.3 Hz), 6.78-6.69 (2H, m), 6.08 (2H, s), 4.75 (1H, d, J = 8.1 Hz), 3.88-3.62 (3H, m), 3.49 (1H, s), 3.09-3.01 (2H, m), 2.97-2.85 (2H, m), 2.84-2.72 (2H, m), 1.84 (1H, septet, J = 6.6 Hz), 1.37 (9H, s), 0.90 (3H, d, J =

6.6 Hz), 0.86 (3H, d, J = 6.6 Hz); coupled LCMS showed the product as a single major peak with m/z 537 $[M+H]^+$ at R_T 2.50 min.

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Example 13



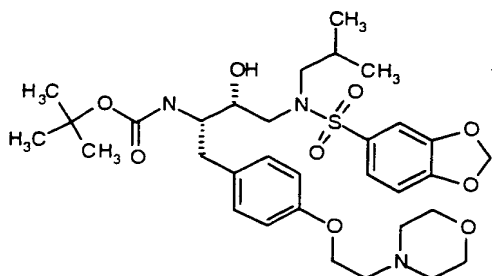
**Boc methylenedioxybenzenesulfonamide acetylmorpholine
tethered product (21)**

5

A solution of Boc methylenedioxybenzenesulfonamide free phenol (0.036 g, 0.067 mmol) in 1,4-dioxane (1 mL) was combined with cesium carbonate (0.055 g, 0.168 mmol) and 4-(2-chloroacetyl)morpholine (0.016 g, 0.100 mmol). The solution was heated to 85 °C with stirring for 2 hours, and was cooled and dried *in vacuo* to give a pale yellow oil. Flash column chromatography (ethyl acetate) gave Boc methylenedioxybenzenesulfonamide acetylmorpholine tethered product (21) (0.032 g, 71%) as a white foam: R_f = 0.51 (ethyl acetate); ^1H NMR (CDCl_3); 7.34 (1H, dd, J = 8.0, 1.7 Hz), 7.22-7.14 (3H, m), 6.93-6.84 (3H, m), 6.09 (2H, s), 4.70-4.59 (1H, m), 4.67 (2H, s), 3.97-3.90 (1H, m), 3.82-3.58 (10H, m), 3.12-3.04 (2H, m), 2.99-2.78 (4H, m), 1.84 (1H, septet, J = 6.6 Hz), 1.36 (9H, s), 0.91 (3H, d, J = 6.6 Hz), 0.88 (3H, d, J = 6.6 Hz); coupled LCMS showed the product as a single major peak with m/z 664 $[\text{M}+\text{H}]^+$; HPLC (205 nm) showed the material as a single major peak at R_T of 2.38 min. (purity = 99%).

25

Example 14

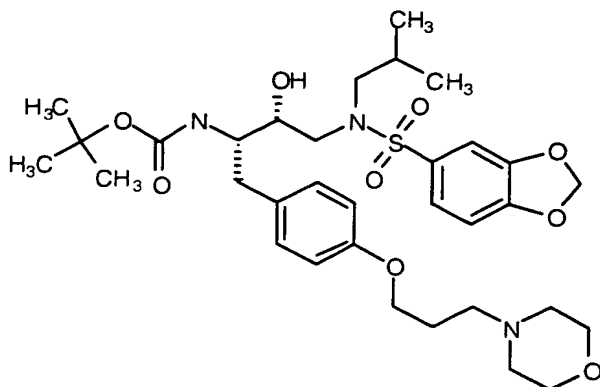


**Boc methylenedioxybenzenesulfonamide ethylmorpholine
5 tethered product (17)**

A solution of Boc methylenedioxybenzenesulfonamide free
phenol (0.034 g, 0.063 mmol) in 1,4-dioxane (1 mL) was
combined with cesium carbonate (0.052 g, 0.158 mmol) and
10 N-(2-chloroethyl)morpholine (0.014 g, 0.095 mmol). The
solution was heated to 85 °C with stirring for 5 hours,
and was cooled and dried in vacuo to give a pale yellow
oil. Flash column chromatography (4:1 ethyl
acetate/hexane) gave Boc methylenedioxybenzenesulfonamide
15 ethylmorpholine tethered product (17) (0.012 g, 29%) as a
pale yellow oil: R_f = 0.32 (4:1 ethyl acetate/hexane); ^1H
NMR (CDCl_3); 7.32 (1H, dd, J = 8.2, 1.5 Hz), 7.18 (1H, d,
 J = 1.5 Hz), 7.15 (2H, d, J = 8.3 Hz), 6.88 (1H, d, J =
8.2 Hz), 6.84 (2H, d, J = 8.3 Hz), 6.09 (2H, s), 4.63
20 (1H, d, J = 8.2 Hz), 3.93 (1H, br. s), 3.83-3.64 (7H, m),
3.06 (2H, d, J = 4.9 Hz), 2.98-2.76 (6H, m), 2.68-2.55
(4H, m), 2.54-2.48 (1H, m), 1.84 (1H, septet, J = 6.7
Hz), 1.35 (9H, s), 0.90 (3H, d, J = 6.7 Hz), 0.87 (3H, d,
 J = 6.7 Hz); coupled LCMS showed the product as a single
25 major peak with m/z 650 $[\text{M}+\text{H}]^+$; HPLC (205 nm) showed the
material as a single major peak at R_T of 2.06 min. (purity
= 92%).

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Example 15



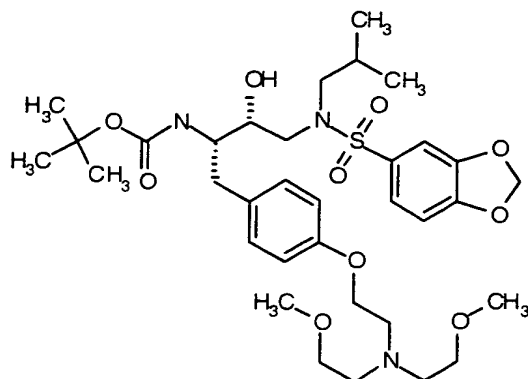
5 **Boc methylenedioxybenzenesulfonamide propylmorpholine
tethered product (20)**

A solution of Boc methylenedioxybenzenesulfonamide free
phenol (0.038 g, 0.071 mmol) in 1,4-dioxane (1 mL) was
10 combined with cesium carbonate (0.058 g, 0.177 mmol) and
N-(3-chloropropyl)morpholine (0.017 g, 0.107 mmol). The
solution was heated to 85 °C with stirring for 8 hours,
and was cooled and dried in vacuo to give a pale yellow
oil. Flash column chromatography (4:1 ethyl
15 acetate/hexane) gave Boc methylenedioxybenzenesulfonamide
propylmorpholine tethered product (20) (0.006 g, 13%) as
a pale yellow oil: R_f = 0.15 (4:1 ethyl acetate/hexane);
 ^1H NMR (CDCl_3); 7.32 (1H, dd, J = 8.1, 1.8 Hz), 7.19 (1H,
d, J = 1.8 Hz), 7.14 (2H, d, J = 8.4 Hz), 6.88 (1H, d, J
20 = 8.1 Hz), 6.83 (2H, d, J = 8.4 Hz), 6.08 (2H, s), 4.62
(1H, br. s), 4.00 (2H, t, J = 6.3 Hz), 3.84-3.65 (8H, m),
3.61 (2H, t, J = 6.6 Hz), 3.10-3.03 (2H, m), 3.00-2.77
(3H, m), 2.64-2.35 (4H, m), 2.05-1.91 (2H, m), 1.85 (1H,
septet, J = 6.6 Hz), 1.37 (9H, s), 0.90 (3H, d, J = 6.6
25 Hz), 0.87 (3H, d, J = 6.6 Hz); coupled LCMS showed the
product as a single major peak with m/z 664 $[\text{M}+\text{H}]^+$; HPLC

(205 nm) showed the material as a single major peak at R_T of 2.23 min. (purity = 80%).

5

Example 16



Boc methylenedioxybenzenesulfonamide bis-methoxyethylamine tethered product (18)

- 10 A solution of Boc methylenedioxybenzenesulfonamide free phenol (0.034 g, 0.063 mmol) in 1,4-dioxane (1 mL) was combined with cesium carbonate (0.052 g, 0.158 mmol) and freshly prepared 2-[N,N-bis-(2-methoxyethyl)amino]ethyl chloride (ca. 0.031 g, ca. 0.016 mmol). The solution was
- 15 heated to 85 °C with stirring for 3.5 hours, and was cooled and dried in vacuo to give a pale yellow oil. Flash column chromatography (ethyl acetate) gave Boc methylenedioxybenzenesulfonamide bis-methoxyethylamine tethered product (18) (0.024 g, 54%) as a white foam: R_f
- 20 = 0.35 (ethyl acetate); ^1H NMR (CDCl_3) 7.32 (1H, dd, J = 8.2, 1.6 Hz), 7.18 (1H, d, J = 1.6 Hz), 7.14 (2H, d, J = 8.5 Hz), 6.88 (1H, d, J = 8.2 Hz), 6.83 (2H, d, J = 8.5 Hz), 6.08 (2H, s), 4.63 (1H, d, J = 7.7 Hz), 4.03 (2H, t, J = 6.1 Hz), 3.92 (1H, br. s), 3.81-3.73 (1H, m), 3.73-
- 25 3.63 (1H, m), 3.50 (4H, t, J = 5.8 Hz), 3.34 (6H, s), 3.10-3.03 (2H, m), 3.01 (2H, t, J = 6.1 Hz), 2.98-2.75

5 showed the material as a single major peak at R_T of 2.20 min. (purity = 100%).

Example 17

10 Boc methylenedioxybenzenesulfonamide 3-picolyyl tethered
product (intermediate en route to compound 53)

A solution of Boc methylenedioxybenzenesulfonamide free phenol (0.010 g, 0.020 mmol) in 1,4-dioxane (1 mL) was combined with cesium carbonate (0.015 g, 0.047 mmol), 3-(chloromethyl)pyridine (ca. 0.004 g, ca. 0.030 mmol; prepared by dissolving 10 mg of 3-(chloromethyl)pyridine hydrochloride in sodium hydroxide (1.5 mL) and diethyl ether (1.5 mL), the organic extract was dried over MgSO_4 and the solvent removed in vacuo) and potassium iodide (~1 mg, 0.006 mmol). The solution was heated to 60 °C with stirring for 8 hours, and was cooled and dried in vacuo to give a pale yellow oil. Flash column chromatography (3:7 ethyl acetate/hexane) gave Boc methylenedioxybenzenesulfonamide 3-picolyl tethered product (0.004 g, 34%) as a colourless oil: R_f = 0.05 (1:1 ethyl acetate/hexane); coupled LCMS showed the product as a single major peak with m/z 628 $[\text{M}+\text{H}]^+$ at R_T of 2.27 min.

Example 18

Boc methylenedioxybenzenesulfonamide 2-picolyl tethered product (intermediate en route to 52)

5 A solution of Boc methylenedioxybenzenesulfonamide free phenol (0.010 g, 0.020 mmol) in 1,4-dioxane (1 mL) was combined with cesium carbonate (0.015 g, 0.047 mmol), 2-(chloromethyl)pyridine (ca. 0.004 g, ca. 0.030 mmol; prepared by dissolving 10 mg of 2-(chloromethyl)pyridine hydrochloride in sodium hydroxide (1.5 mL) and diethyl
10 ether (1.5 mL), the organic extract was dried over MgSO_4 and the solvent removed *in vacuo*) and potassium iodide (~1 mg, 0.006 mmol). The solution was heated to 60 °C with stirring for 8 hours, and was cooled and dried *in vacuo* to give a pale yellow oil. Flash column chromatography
15 (3:7 ethyl acetate/hexane) gave Boc methylenedioxybenzenesulfonamide 2-picolyl tethered product (0.004 g, 34%) as a colourless oil: R_f = 0.3 (1:1 ethyl acetate/hexane); coupled LCMS showed the product with m/z 628 $[\text{M}+\text{H}]^+$ at R_T of 2.35 min.

20

Example 19

Boc methylenedioxybenzenesulfonamide 4-picolyl tethered product (intermediate en route to compound 54)

25 A solution of Boc methylenedioxybenzenesulfonamide free phenol (0.010 g, 0.020 mmol) in 1,4-dioxane (1 mL) was combined with cesium carbonate (0.015 g, 0.047 mmol), 4-(chloromethyl)pyridine (ca. 0.004 g, ca. 0.030 mmol; prepared by dissolving 10 mg of 4-(chloromethyl)pyridine
30 hydrochloride in sodium hydroxide (1.5 mL) and diethyl ether (1.5 mL), the organic extract was dried over MgSO_4 and the solvent removed *in vacuo*) and potassium iodide (~1 mg, 0.006 mmol). The solution was heated to 60 °C with

stirring for 16 hours, and was cooled and dried in vacuo to give a pale yellow oil. Flash column chromatography (3:7 ethyl acetate/hexane) gave Boc methylenedioxybenzenesulfonamide 4-picolyl tethered product (0.004 g, 34%) as a colourless oil: R_f = 0.10 (2:3 ethyl acetate/hexane); coupled LCMS showed the product with m/z 628 $[M+H]^+$ at R_T of 2.24 min.

Example 20

10 **Boc methylenedioxybenzenesulfonamide 3-methyl-5-methylisoxazole tethered product**
(intermediate en route to compound 55)

005090" 1976560
15 A solution of Boc methylenedioxybenzenesulfonamide free phenol (0.010 g, 0.020 mmol) in 1,4-dioxane (1 mL) was combined with cesium carbonate (0.015 g, 0.047 mmol), 3-chloromethyl-5-methyl isoxazole (0.004 g, 0.028 mmol) and potassium iodide (~1 mg, 0.006 mmol). The solution was heated to 60 °C with stirring for 16 hours, and was cooled
20 and dried in vacuo to give a pale yellow oil. Flash column chromatography (3:7 ethyl acetate/hexane) gave Boc methylenedioxybenzenesulfonamide 3-methyl-5-methylisoxazole tethered product (0.005 g, 42%) as a colourless oil: R_f = 0.25 (3:7 ethyl acetate/hexane); 1H
25 NMR ($CDCl_3$) 7.34 (1H, dd, J = 8.4, 1.8 Hz), 7.21-7.13 (3H, m), 6.90 (3H, m), 6.11 (1H, s), 6.09 (2H, s), 5.09 (2H, s), 4.65 (1H, d, J = 8.2 Hz), 3.92 (1H, br. s), 3.82-3.75 (1H, m), 3.73-3.63 (1H, m), 3.09-3.03 (2H, m), 2.98-2.80 (4H, m), 2.43 (3H, s), 1.84 (1H, septet, J = 6.6 Hz),
30 1.36 (9H, s), 0.91 (3H, d, J = 6.6 Hz), 0.88 (3H, d, J = 6.6 Hz).

Example 21A

**Boc methylenedioxybenzenesulfonamide 1-methyl-3,5-
dimethylpyrazole tethered product
(intermediate en route to compound 56)**

5 A solution of Boc methylenedioxybenzenesulfonamide free
phenol (0.010 g, 0.020 mmol) in DMF (1 mL) was combined
with anhydrous cesium carbonate (0.015 g, 0.047 mmol).
Freshly prepared 1-chloromethyl-3,5-dimethylpyrazole
hydrochloride (0.006 g, 0.033 mmol) in DMF (0.5 ml) was
10 added dropwise. The solution was heated to 60 °C with
stirring for 15 minutes, and was cooled and dried in
vacuo to give a green oil. Flash column chromatography
(1:1 ethyl acetate/hexane) gave Boc
methylenedioxybenzenesulfonamide 1-methyl-3,5-
15 dimethylpyrazole tethered product (0.002 g, 17%) as a
colourless oil: R_f = 0.25 (1:1 ethyl acetate/hexane);
coupled LCMS showed the product as a single major peak
with m/z 645 $[M+H]^+$ at R_T of 1.68 min.

20

Example 21B

**Boc methylenedioxybenzenesulfonamide ethylpyrazole
tethered product (intermediate en route to compound 57)**

A solution of Boc methylenedioxybenzenesulfonamide free
25 phenol (0.010 g, 0.020 mmol) in acetone (1 mL) was
combined with cesium carbonate (0.015 g, 0.047 mmol), 1-
(2-chloroethyl)pyrazole (0.004 g, 0.031 mmol) and sodium
iodide (~1 mg, 0.007 mmol). The solution was heated to
55 °C with stirring for 60 hours, and was cooled and dried
30 in vacuo to give a yellow oil. Flash column
chromatography (1:3 ethyl acetate/hexane) gave Boc
methylenedioxybenzenesulfonamide ethylpyrazole tethered
product (0.0015 g, 13%) as a colourless oil: R_f = 0.20

5

Example 22

Bis-THF *m*-nitrobenzenesulfonamide Benzyl Ether
(intermediate en route to compounds:25, 34-37)

5

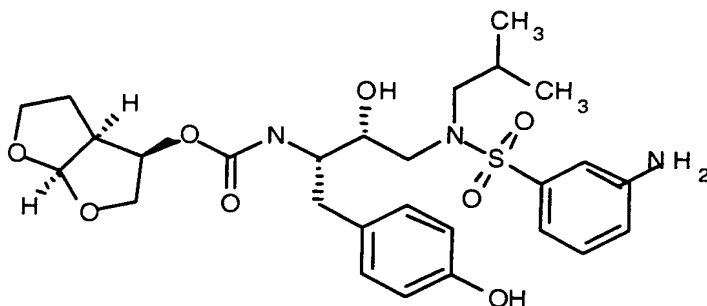
Boc *m*-nitrobenzenesulfonamide benzyl (5c; 0.050 g, 0.08 mmol) was dissolved in DCM (3.4 mL). Trifluoroacetic acid (1.6 mL) was added dropwise, and the reaction was stirred at room temperature for 1 hour. The solvent was removed in vacuo, and the resultant orange oil was dissolved in DCM (1.5 mL) and the solution cooled to 0°C using an ice bath. Diisopropylethylamine (0.42 mL, 2.39 mmol) was added dropwise with stirring, followed by (3R, 3aS, 6aR) hexahydrofuro[2,3-*b*]furan-2-yl 4-nitrophenyl carbonate (0.026 g, 0.088 mmol) in one portion as a solid. After 5 minutes, the ice bath was removed and the reaction mixture was allowed to warm to room temperature with stirring overnight. The solvent was removed in vacuo, and the resultant yellow oil was purified by flash column chromatography (1:1 ethyl acetate/hexane) to give bis-THF *m*-nitrobenzenesulfonamide benzyl ether (6; 0.036 g, 66%) as a white foam:

R_f = 0.39 (1:1 ethyl acetate/hexane); ^1H NMR (CDCl_3) 8.62 (1H, d, J = 2.3 Hz), 8.42 (1H, dt, J = 8.1, 0.9 Hz), 8.10 (1H, d, J = 7.7 Hz), 7.73 (1H, t, J = 8.1 Hz), 7.45-7.29 (5H, m), 7.12 (2H, d, J = 8.6 Hz), 6.89 (2H, d, J = 8.6 Hz), 5.65 (1H, d, J = 5.4 Hz), 5.10- 4.98 (1H, m), 5.02 (2H, s), 4.93 (1H, d, J = 8.1 Hz), 3.96 (1H, dd, J = 9.0, 6.3 Hz), 3.91-3.76 (3H, m), 3.76-3.61 (2H, m), 3.24 (1H, dd, J = 15.4, 8.1 Hz), 3.16 (1H, dd, 15.2, 3.0 Hz), 3.06-2.96 (3H, m), 2.96-2.88 (1H, m), 2.74 (1H, dd, J = 14.2, 9.3 Hz), 1.88 (1H, septet, J = 6.7 Hz), 1.73-1.59

(1H, m), 1.59-1.48 (1H, m), 0.90 (6H, t, J = 6.3 Hz), OH signal not observed.

Example 23

5



Bis-THF m-aminophenylsulfonamide Benzyl Ether and Bis-THF m-aminophenylsulfonamide Free Phenol (intermediates en route to compounds 34-37)

10

A solution of *bis*-THF *m*-nitrobenzenesulfonamide benzyl ether (5c; 0.036 g, 0.053 mmol) and 10% palladium on carbon (wet Degussa) (0.012 g) in degassed ethyl acetate (15 mL) was stirred under a balloon of hydrogen. The reaction was monitored by TLC, and after 2 hours the starting material had been consumed. An aliquot (5 mL) of the reaction mixture was removed, and filtered through a pad of celite. The filtrate was dried *in vacuo* to give *bis*-THF *m*-aminophenylsulfonamide benzyl ether (compound 22) as an off-white oily solid (0.010 g, 29%).

15

20

A fresh portion of 10% palladium on carbon (wet Degussa) (0.012 g) was added to the remainder of the reaction mixture, and the reaction stirred under a balloon of hydrogen for a further 2 hours. TLC indicated that an amount of intermediate product remained, so a further portion of 10% palladium on carbon (wet Degussa) (0.012 g) was added to the reaction mixture, and the

25

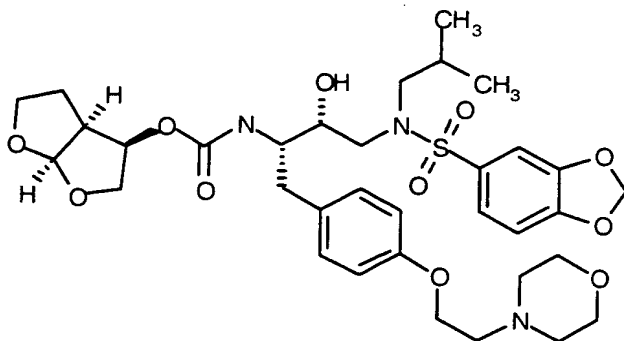
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reaction stirred under a balloon of hydrogen overnight.
The reaction mixture was filtered through a pad of
celite, and the filtrate was dried *in vacuo* to give *bis*-
THF *m*-aminophenylsulfonamide free phenol (6: R8 = H, D'=
5 *i*-Bu, E = *m*-NH₂Ph, A = *bis*-THF- CO-) as an off-white oily
solid (0.016g, 54%).

For *bis*-THF *m*-aminophenylsulfonamide benzyl ether: R_f =
0.47 (3:1 ethyl acetate/hexane); ¹H NMR (CDCl₃); 7.45-7.28
10 (7H, m), 7.28-7.19 (1H, m), 7.17-7.08 (3H, m), 6.89 (2H,
d, J = 8.1 Hz), 5.65 (1H, dd, J = 9.3, 5.2 Hz), 5.10-4.95
(4H, m), 4.00-3.60 (6H, m), 3.30-2.80 (6H, m), 2.74 (1H,
dd, J = 10.4, 9.0 Hz), 1.92-1.80 (1H, m), 1.75-1.43 (2H,
m), 0.95-0.88 (6H, m); OH and NH₂ signals not observed;
15 coupled LCMS showed the product as a single major peak
with *m/z* 654 [M+H]⁺; HPLC (205 nm) showed the material to
be split into 2 peaks at R_Ts of 2.33 min. & 2.38 min.
(combined purity = 89%).

20 For *bis*-THF *m*-aminophenylsulfonamide free phenol : D' = *i*-
Bu, E = *m*-NH₂Ph, A = *bis*-THF- CO-, R8 = H): R_f = 0.21
(3:1 ethyl acetate/hexane); ¹H NMR (CDCl₃); 7.28 (1H, dd,
J = 15.3, 7.2 Hz), 7.13-7.10 (1H, m), 7.08 (2H, d, J =
8.1 Hz), 7.03-6.99 (1H, m), 6.86 (1H, dd, J = 7.9, 1.6
25 Hz), 6.75 (2H, d, J = 8.6 Hz), 5.66 (1H, d, J = 4.9 Hz),
5.05 (2H, d, J = 7.7 Hz), 4.01- 3.59 (7H, m), 3.17-3.04
(1H, m), 3.04-2.89 (4H, m), 2.86-2.74 (2H, m), 1.91-1.50
(3H, m), 0.93 (3H, d, J = 6.5 Hz), 0.81 (3H, d, J = 6.5
Hz); OH and NH₂ signals not observed; coupled LCMS showed
30 the product as a single major peak with *m/z* 565 [M+H]⁺;
HPLC (205 nm) showed the material as a single major peak
at R_T of 1.53 min. (purity = 92%).

Example 25



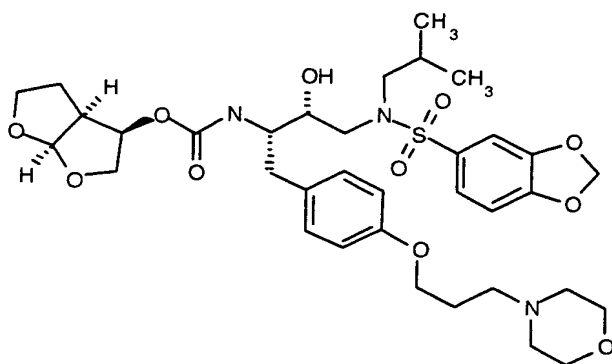
**Bis-THF methylenedioxybenzenesulfonamide
ethylmorpholine tethered product (Compound 48)**

5

Boc methylenedioxybenzenesulfonamide ethylmorpholine
tethered product (Compound 17; 0.011 g, 0.017 mmol) was
dissolved in DCM (1 mL). Trifluoroacetic acid (0.5 mL)
was added dropwise, and the reaction was stirred at room
10 temperature for 1 hour. The solvent was removed in
vacuo, and the resultant orange oil was dissolved in DCM
(0.3 mL) and the solution cooled to 0°C using an ice bath.
Diisopropylethylamine (0.088 mL, 0.51 mmol) was added
dropwise with stirring, followed by (3R, 3aS, 6aR)
15 hexahydrofuro[2,3-b]furan-2-yl 4-nitrophenyl carbonate
(0.006 g, 0.020 mmol) in one portion as a solid. After 5
minutes, the ice bath was removed and the reaction
mixture was allowed to warm to room temperature with
stirring overnight. The solvent was removed in vacuo,
20 and the resultant yellow oil was purified by flash column
chromatography (from 4:1 ethyl acetate/hexane to 9:1
DCM/methanol) to give bis-THF
methylenedioxybenzenesulfonamide ethylmorpholine tethered
product (Compound 48) (0.0095 g, 79%) as a pale yellow
25 oil: R_f = 0.50 (9:1 DCM/methanol); ^1H NMR (CDCl_3) 7.40-
7.28 (1H, m), 7.23-7.04 (3H, m), 6.97-6.87 (1H, m), 6.87-
6.77 (2H, m), 6.12 (2H, s), 5.68 (1H, d, J = 4.8 Hz),

5.11-5.00 (1H, m), 5.00-4.87 (1H, m), 4.03-3.92 (2H, m),
 3.92-3.53 (10H, m), 3.21-3.07 (2H, m), 3.07-2.92 (3H, m),
 2.92-2.74 (4H, m), 2.74-2.54 (4H, m), 1.96-1.37 (3H, m),
 1.02-0.78 (6H, m), OH signal not observed; coupled LCMS
 5 showed the product as a single major peak with m/z 706
 [M+H]⁺; HPLC (205 nm) showed the material as a single
 major peak at R_T of 1.83 min. (purity = 95%).

Example 26



**Bis-THF methylenedioxybenzenesulfonamide
 propylmorpholine tethered product (Compound 49)**

Boc methylenedioxybenzenesulfonamide
 15 propylmorpholine tethered product (20) (0.006 g, 0.009
 mmol) was dissolved in DCM (1 mL). Trifluoroacetic acid
 (0.5 mL) was added dropwise, and the reaction was stirred
 at room temperature for 1 hour. The solvent was removed
 in vacuo, and the resultant orange oil was dissolved in
 20 DCM (0.3 mL) and the solution cooled to 0°C using an ice
 bath. Diisopropylethylamine (0.047 mL, 0.27 mmol) was
 added dropwise with stirring, followed by (3R, 3aS, 6aR)
 hexahydrofuro[2,3-b]furan-2-yl 4-nitrophenyl carbonate
 (0.003 g, 0.011 mmol) in one portion as a solid. After 5
 25 minutes, the ice bath was removed and the reaction
 mixture was allowed to warm to room temperature with

stirring overnight. The solvent was removed *in vacuo*, and the resultant yellow oil was purified by flash column chromatography (from 4:1 ethyl acetate/hexane to 9:1 DCM/methanol) to give *bis*-THF

5 methylenedioxybenzenesulfonamide propylmorpholine
tethered product (Compound 49) (0.0045 g, 69%) as a
yellow oil: R_f = 0.50 (9:1 DCM/methanol); ^1H NMR (CDCl_3)
7.36-7.30 (1H, m), 7.19 (1H, s), 7.13 (2H, d, J = 8.4
Hz), 6.91 (1H, d, J = 8.3 Hz), 6.82 (2H, d, J = 8.3 Hz),
10 6.11 (2H, s), 5.67 (1H, d, J = 5.1 Hz), 5.19-5.08 (1H,
dd, J = 13.6, 6.1 Hz), 4.92 (1H, d, J = 9.0 Hz), 4.07-
3.92 (4H, m), 3.92-3.65 (8H, m), 3.19-3.08 (1H, m), 3.05-
2.85 (4H, m), 2.85-2.71 (2H, m), 2.71-2.43 (6H, m), 2.13-
1.94 (2H, m), 1.94-1.77 (1H, m), 1.77-1.49 (2H, m), 0.95
15 (3H, d, J = 6.5 Hz), 0.90 (3H, d, J = 6.7 Hz), OH signal
not observed; coupled LCMS showed the product as a single
major peak with m/z 720 $[\text{M}+\text{H}]^+$; HPLC (205 nm) showed the
material as a single major peak at R_T of 1.90 min. (purity
= 93%).

20

Example 27

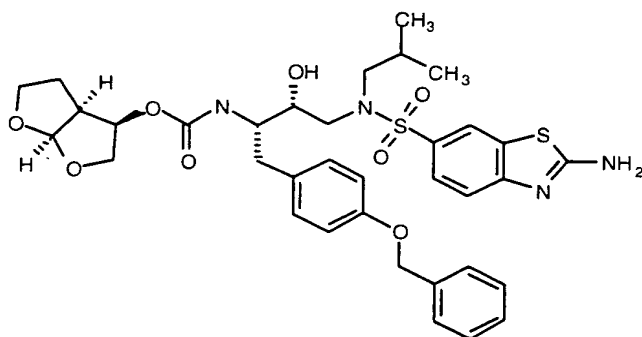
Boc Benzothiazole Sulfonamide Benzyl Ether (Compound 5g, Scheme 1)

25

The isobutylamino alcohol (0.50 g, 1.13 mmol) was
combined with 2-aminobenzothiazole-6- sulfonyl chloride
(0.373 g, 1.50 mmol) in anhydrous DCM (10 mL) under a N_2
atmosphere. Diisopropylethylamine (0.59 mL, 3.39 mmol)
30 was added dropwise, and the reaction was stirred at room
temperature for 42 hours. The solvent was removed *in*
vacuo, and the resultant yellow oil was purified by flash
column chromatography (4:1 ethyl acetate/hexane) to give

Boc benzothiazole sulfonamide benzyl ether (5: D' = i-Bu, E = 2-aminobenzothiazole) (0.30 g, 42%) as a white solid: R_f = 0.60 (4:1 ethyl acetate/hexane); ^1H NMR (CDCl_3) 8.05 (1H, d, J = 1.8 Hz), 7.72 (1H, dd, J = 8.6, 1.8 Hz), 7.60 (1H, d, J = 8.6 Hz), 7.49-7.32 (5H, m), 7.19 (2H, d, J = 8.6 Hz), 6.94 (2H, d, J = 8.6 Hz), 5.67 (2H, br.s, NH_2), 5.07 (2H, s), 4.70 (1H, br.d, J = 7.7 Hz), 4.00 (1H, br.s, OH), 3.89-3.80 (1H, m), 3.80-3.66 (1H, m), 3.23-3.06 (2H, m), 3.06-2.77 (4H, m), 1.89 (1H, septet, J = 7.2 Hz), 1.38 (9H, s), 0.94 (3H, d, J = 6.3 Hz), 0.90 (3H, d, J = 6.3 Hz).

Example 28

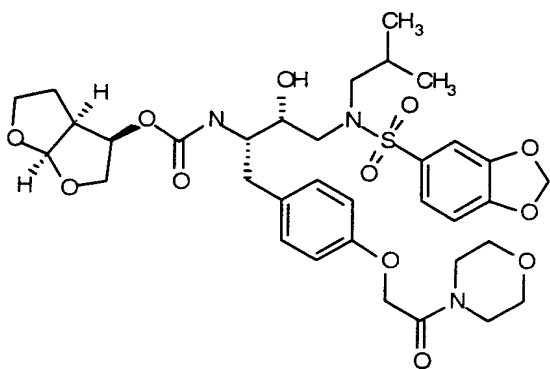


**Bis-THF benzothiazolesulfonamide
Benzyl Ether (Compound 44)**

Boc benzothiazole sulfonamide benzyl ether (Compound 5, Scheme 1: D' = i-Bu, E = 2-aminobenzothiazole) (0.30 g, 0.46 mmol) was dissolved in DCM (10 mL). Trifluoroacetic acid (5 mL) was added dropwise, and the reaction was stirred at room temperature for 1 hour. The solvent was removed *in vacuo*, and the resultant orange oil was dissolved in DCM (15 mL). Triethylamine (1.28 mL, 9.20 mmol) was added dropwise with stirring, followed by (3R, 3aS, 6aR) hexahydrofuro[2,3-*b*]furan-2-yl 4-nitrophenyl

carbonate (0.149 g, 0.50 mmol) in one portion as a solid. The reaction mixture was allowed to stir at room temperature overnight. The solvent was removed in vacuo, and the resultant yellow oil was purified by flash column chromatography (ethyl acetate) to give bis-THF benzothiazole sulfonamide benzyl ether (Compound 44) (0.158 g, 48%) as a white solid: R_f = 0.50 (ethyl acetate); ^1H NMR (CDCl_3) 8.03 (1H, s), 7.71 (1H, d, J = 8.6 Hz), 7.58 (1H, d, J = 8.6 Hz), 7.48-7.32 (5H, m), 7.15 (2H, d, J = 8.1 Hz), 6.91 (2H, d, J = 8.6 Hz), 5.93 (2H, br.s, NH_2), 5.67 (1H, d, J = 5.4 Hz), 5.16 (1H, d, J = 8.6 Hz), 5.04 (2H, s), 4.05-3.60 (7H, m), 3.27-3.15 (1H, m), 3.12-2.97 (4H, m), 2.97-2.83 (2H, m), 2.78 (1H, dd, J = 14.3, 8.8 Hz), 1.95-1.48 (3H, m), 0.96 (3H, d, J = 6.8 Hz), 0.92 (3H, d, J = 6.8 Hz); coupled LCMS showed the product as a single major peak with m/z 712 $[\text{M}+\text{H}]^+$; HPLC (205 nm) showed the material as a single major peak at R_T of 2.26 min. (purity = 100%).

Example 29

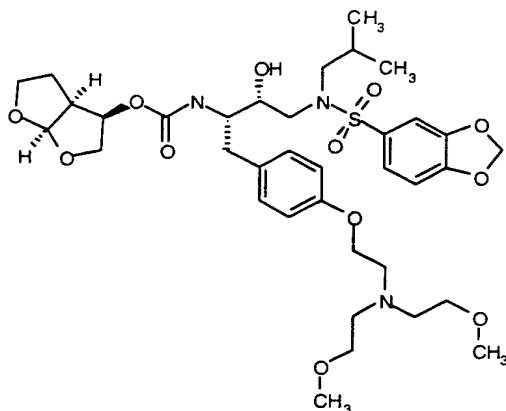


Bis-THF methylenedioxybenzenesulfonamide acetylmorpholine tethered product (50)

Boc methylenedioxybenzenesulfonamide acetylmorpholine tethered product (21) (0.027 g, 0.041 mmol) was dissolved

in DCM (2 mL). Trifluoroacetic acid (1 mL) was added dropwise, and the reaction was stirred at room temperature for 1 hour. The solvent was removed *in vacuo*, and the resultant orange oil was dissolved in DCM (1.5 mL) and the solution cooled to 0 °C using an ice bath. Diisopropylethylamine (0.21 mL, 1.22 mmol) was added dropwise with stirring, followed by (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-2-yl 4-nitrophenyl carbonate (0.013 g, 0.045 mmol) in one portion as a solid. After 5 minutes, the ice bath was removed and the reaction mixture was allowed to warm to room temperature with stirring overnight. The solvent was removed *in vacuo*, and the resultant yellow oil was purified by flash column chromatography (ethyl acetate to methanol) to give *bis*-THF methylenedioxybenzenesulfonamide acetylmorpholine tethered product (50) (0.011 g, 38%) as a yellow foam: R_f = 0.10 (ethyl acetate); ^1H NMR (CDCl_3) 7.33 (1H, d, J = 8.1 Hz), 7.20-7.08 (3H, m), 6.94-6.83 (3H, m), 6.09 (2H, s), 5.72-5.61 (1H, m), 5.10-4.97 (2H, m), 4.67 (2H, s), 4.02-3.92 (1H, m), 3.91-3.77 (3H, m), 3.76-3.56 (10H, m), 3.19-3.07 (1H, m), 3.06-2.88 (4H, m), 2.87-2.71 (2H, m), 1.89-1.75 (1H, m), 1.74-1.62 (1H, m), 1.61-1.48 (1H, m), 0.93 (3H, d, J = 6.6 Hz), 0.89 (3H, d, J = 6.6 Hz), OH signal not observed; coupled LCMS showed the product as a single major peak with m/z 720 $[\text{M}+\text{H}]^+$; HPLC (205 nm) showed the material as a single major peak at R_T of 2.02 min. (purity = 88%).

Example 32

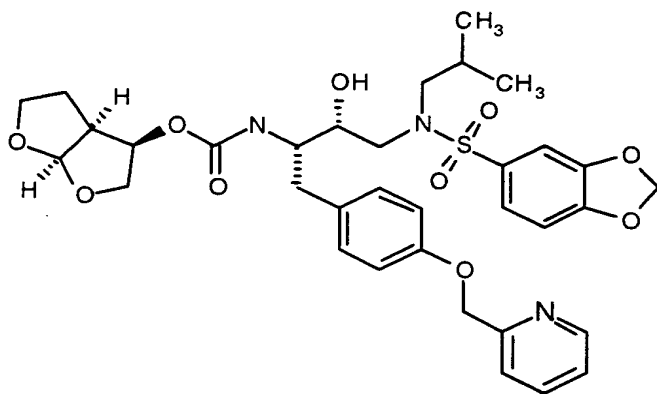


Bis-THF methylenedioxybenzenesulfonamide bis-methoxyethylamine tethered product (51)

5 Boc methylenedioxybenzenesulfonamide bis-methoxyethylamine tethered product (18) (0.023 g, 0.033 mmol) was dissolved in DCM (1 mL). Trifluoroacetic acid (0.5 mL) was added dropwise, and the reaction was stirred at room temperature for 1 hour. The solvent was removed
10 in vacuo, and the resultant orange oil was dissolved in DCM (1 mL) and the solution cooled to 0 °C using an ice bath. Diisopropylethylamine (0.173 mL, 0.99 mmol) was added dropwise with stirring, followed by (3R, 3aS, 6aR) hexahydrofuro[2,3-b]furan-2-yl 4-nitrophenyl carbonate
15 (0.012 g, 0.040 mmol) in one portion as a solid. After 5 minutes, the ice bath was removed and the reaction mixture was allowed to warm to room temperature with stirring overnight. The solvent was removed in vacuo, and the resultant yellow oil was purified by flash column
20 chromatography (4:1 ethyl acetate/hexane to ethyl acetate to 9:1 DCM/methanol) to give bis-THF methylenedioxybenzenesulfonamide bis-methoxyethylamine tethered product (51) (0.022 g, 89%) as a yellow foam: R_f = 0.50 (9:1 DCM/methanol); ^1H NMR (CDCl_3) 7.33 (1H, dd, J = 8.2, 1.8 Hz), 7.17 (1H, d, J = 1.8 Hz), 7.13 (2H, d, J

= 8.6 Hz), 6.89 (1H, d, J = 8.2 Hz), 6.80 (2H, d, J = 8.6 Hz), 6.09 (2H, s), 5.65 (1H, d, J = 5.2 Hz), 5.08-4.98 (2H, m), 4.35-4.24 (2H, m), 4.00-3.90 (1H, m), 3.90-3.63 (6H, m), 3.61-3.49 (2H, m), 3.42-3.31 (2H, m), 3.35 (6H, s), 3.16-3.07 (2H, m), 3.05-2.88 (4H, m), 2.88-2.74 (2H, m), 1.83 (1H, septet, J = 6.6 Hz), 1.74-1.54 (1H, m), 1.48-1.35 (5H, m), 0.92 (3H, d, J = 6.6 Hz), 0.88 (3H, d, J = 6.6 Hz), OH signal not observed; coupled LCMS showed the product as a single major peak with m/z 752 $[M+H]^+$; HPLC (205 nm) showed the material as a single major peak at R_T of 1.92 min. (purity = 91%).

Example 33



15

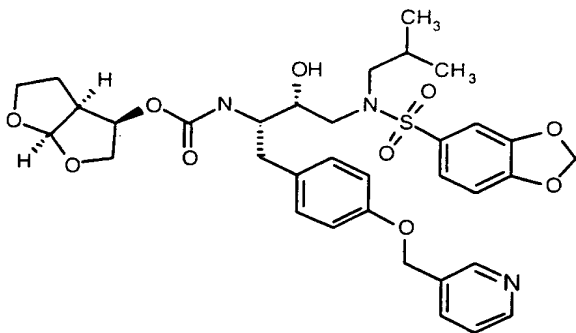
Bis-THF methylenedioxybenzenesulfonamide 2-picolyt tethered product (53)

Boc methylenedioxybenzenesulfonamide.2-picolyt tethered product (0.004 g, 0.006 mmol) was dissolved in DCM (0.6 mL). Trifluoroacetic acid (0.3 mL) was added dropwise, and the reaction was stirred at room temperature for 45 minutes. The solvent was removed *in vacuo*, and the resultant orange oil was dissolved in DCM (1.5 mL). Diisopropylethylamine (0.060 mL, 0.33 mmol) was added dropwise with stirring at room temperature, followed by

(3R, 3aS, 6aR) hexahydrofuro[2,3-b]furan-2-yl 4-nitrophenyl carbonate (0.005 g, 0.017 mmol) in one portion as a solid. The yellow reaction mixture was stirred at room temperature overnight. The solvent was removed *in vacuo*, and the resultant yellow oil was purified by flash column chromatography (from 2:3 ethyl acetate/hexane to 5:1 ethyl acetate/methanol) to give *bis*-THF methylenedioxybenzenesulfonamide 2-picolyt tethered product (53) (0.0013 g, 30%) as a pale yellow oil: $R_f = 0.35$ (5:1 ethyl acetate/methanol); ^1H NMR (CDCl_3) 8.62-8.57 (1H, m), 7.75-7.70 (1H, m), 7.55-7.48 (1H, m), 7.34 (1H, dd, $J = 8.1, 1.8$ Hz), 7.28-7.20 (1H, obscured m), 7.17-7.10 (3H, m), 6.93-6.87 (3H, m), 6.08 (2H, s), 5.65 (1H, d, $J = 5.0$ Hz), 5.17 (2H, br. s), 5.07-5.01 (1H, m), 4.95-4.89 (1H, m), 4.01-3.93 (1H, m), 3.89-3.80 (3H, m), 3.76-3.67 (1H, m), 3.62-3.58 (1H, m), 3.18-3.09 (1H, m), 3.04-2.87 (4H, m), 2.83-2.73 (2H, m), 1.88-1.79 (1H, m), 1.70-1.47 (2H, obscured m), 0.93 (3H, d, $J = 6.3$ Hz), 0.89 (3H, d, $J = 6.3$ Hz), OH signal not observed; coupled LCMS showed the product as a single major peak with m/z 684 $[\text{M}+\text{H}]^+$; integrated LCMS (205 nm) showed the material as a single major peak at R_T of 2.15 min. (purity = 93%).

25

Example 34

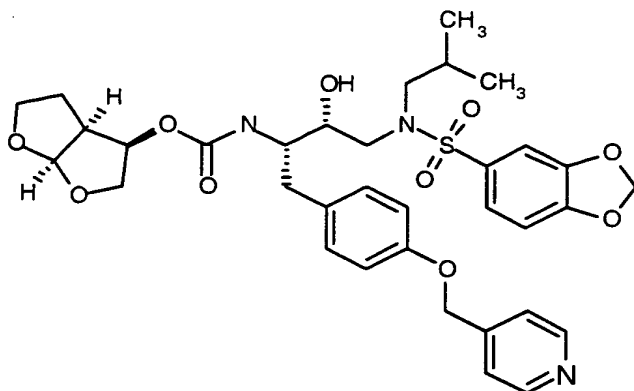


**Bis-THF methylenedioxybenzenesulfonamide 3-picolyl
tethered product (52)**

Boc methylenedioxybenzenesulfonamide 3-picolyl tethered
5 product (0.004 g, 0.006 mmol) was dissolved in DCM (0.6
mL). Trifluoroacetic acid (0.3 mL) was added dropwise,
and the reaction was stirred at room temperature for 45
minutes. The solvent was removed *in vacuo*, and the
resultant orange oil was dissolved in DCM (1 mL).
10 Diisopropylethylamine (0.030 mL, 0.19 mmol) was added
dropwise with stirring at room temperature, followed by
(3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-2-yl 4-
nitrophenyl carbonate (0.003 g, 0.009 mmol) in one
portion as a solid. The yellow reaction mixture was
15 stirred at room temperature overnight. The solvent was
removed *in vacuo*, and the resultant yellow oil was
purified by flash column chromatography (2:3 ethyl
acetate/hexane) to give *bis*-THF
methylenedioxybenzenesulfonamide 3-picolyl tethered
20 product (52) as a pale yellow oil (0.0012 g; an
inseparable mixture with alkylated oxazolidinone): R_f =
0.20 (ethyl acetate); ^1H NMR (CDCl_3) 8.68 (1H, br. s),
8.63-8.56 (1H, m), 7.81-7.74 (1H, m), 7.39-7.30 (1H,
obscured m), 7.34 (1H, dd, J = 8.2, 1.8 Hz), 7.22-7.12
25 (3H, m), 6.94-6.85 (1H, obscured m), 6.90 (2H, d, J = 8.6
Hz), 6.09 (2H, s), 5.67 (1H, d, J = 5.5 Hz), 5.05 (2H,
br. s), 4.95 (1H, d, J = 8.6 Hz), 4.03-3.92 (1H, m),
3.91-3.79 (3H, m), 3.76-3.67 (2H, m), 3.67-3.63 (1H, m),
3.24-3.07 (2H, m), 3.05-2.85 (4H, m), 2.84-2.76 (1H, m),
30 1.91-1.78 (1H, m), 1.77-1.46 (2H, obscured m) 0.91-0.85
(6H, m), OH signal not observed; coupled LCMS showed the
products (alkylated *bis*-THF derivative/alkylated
oxazolidinone) as a single major peak with m/z 684 $[\text{M}+\text{H}]^+$;

5

Example 35



**Bis-THF methylenedioxybenzenesulfonamide 4-picolyl
tethered product (54)**

5

Boc methylenedioxybenzenesulfonamide 4-picolyl tethered product (0.004 g, 0.006 mmol) was dissolved in DCM (0.3 mL). Trifluoroacetic acid (0.1 mL) was added dropwise, and the reaction was stirred at room temperature for 45 minutes. The solvent was removed *in vacuo*, and the resultant orange oil was dissolved in DCM (1 mL). Diisopropylethylamine (0.030 mL, 0.19 mmol) was added dropwise with stirring at room temperature, followed by (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-2-yl 4-nitrophenyl carbonate (0.003 g, 0.009 mmol) in one portion as a solid. The yellow reaction mixture was stirred for 7 hours at room temperature. The solvent was removed *in vacuo*, and the resultant yellow oil was purified by flash column chromatography (2:3 ethyl acetate/hexane to 5:1 ethyl acetate/methanol) to give *bis*-THF methylenedioxybenzenesulfonamide 4-picolyl tethered product (54) as a pale yellow oil (0.0015 g; an inseparable mixture with alkylated oxazolidinone): R_f = 0.40 (5:1 ethyl acetate/methanol); coupled LCMS showed the products (alkylated *bis*-THF derivative/alkylated oxazolidinone) with m/z 684 $[M+H]^+$; HPLC (205 nm) showed

10

15

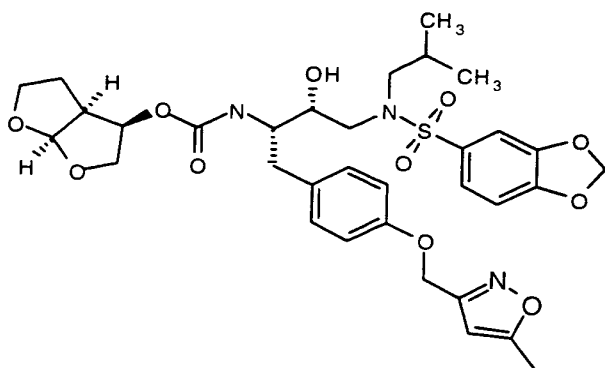
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the materials (alkylated *bis*-THF derivative/alkylated oxazolidinone) as a peak at R_T of 1.97 min. (combined purity = 60%; desired material and alkylated oxazolidinone).

5

Example 36



***Bis*-THF methylenedioxybenzenesulfonamide 3-methyl-5-methylisoxazole tethered product (55)**

10

Boc methylenedioxybenzenesulfonamide 3-methyl-5-methylisoxazole tethered product (0.005 g, 0.008 mmol) was dissolved in DCM (0.3 mL). Trifluoroacetic acid (0.1 mL) was added dropwise, and the reaction was stirred at room temperature for 45 minutes. The solvent was removed *in vacuo*, and the resultant orange oil was dissolved in DCM (1.5 mL). Diisopropylethylamine (0.040 mL, 0.24 mmol) was added dropwise with stirring at room temperature, followed by (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-2-yl 4-nitrophenyl carbonate (0.004 g, 0.012 mmol) in one portion as a solid. The yellow reaction mixture was stirred for 9 hours at room temperature. The solvent was removed *in vacuo*, and the resultant yellow oil was purified by flash column chromatography (from ethyl acetate to 5:1 ethyl acetate/methanol) to give *bis*-THF methylenedioxybenzenesulfonamide 3-methyl-5-

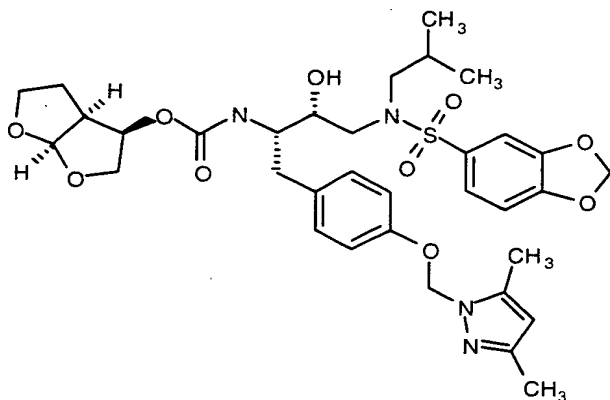
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methylisoxazole tethered product (55) (0.005 g, 97%) as a pale yellow oil: R_f = 0.50 (5:1 ethyl acetate/methanol); ^1H NMR (CDCl_3) 7.33 (1H, dd, J = 8.2, 1.4 Hz), 7.17-7.10 (3H, m), 6.91-6.86 (3H, m), 6.09 (3H, br. s), 5.65 (1H, d, J = 5.0 Hz), 5.06 (2H, s), 4.94 (1H, d, J = 8.7 Hz), 4.00-3.92 (2H, m), 3.88-3.79 (3H, m), 3.74-3.66 (2H, m), 3.64-3.58 (1H, m), 3.17-3.07 (1H, m), 3.03-2.87 (4H, m), 2.82-2.71 (2H, m), 2.42 (3H, s), 1.86-1.76 (1H, m), 1.70-1.49 (2H, m), 0.93 (3H, d, J = 6.6 Hz), 0.88 (3H, d, J = 6.6 Hz); coupled LCMS showed the product as a single major peak with m/z 688 $[\text{M}+\text{H}]^+$; HPLC (205 nm) showed the material as a single major peak at R_T of 2.38 min. (purity = 82%).

Example 37



Bis-THF methylenedioxybenzenesulfonamide 1-methyl-3,5-dimethylpyrazole tethered product (56)

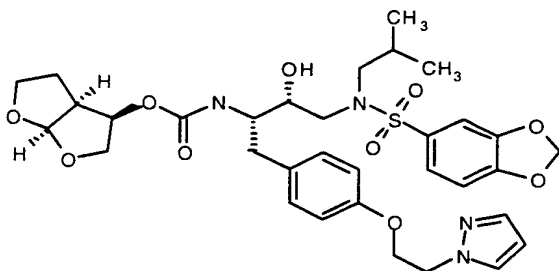
Boc methylenedioxybenzenesulfonamide 1-methyl-3,5-dimethylpyrazole tethered product (0.003 g, 0.005 mmol) was dissolved in DCM (0.15 mL). Trifluoroacetic acid (0.05 mL) was added dropwise, and the reaction was stirred at room temperature for 45 minutes. The solvent was removed *in vacuo*, and the resultant orange oil was dissolved in DCM (1 mL). Diisopropylethylamine (0.030 mL,

0.18 mmol) was added dropwise with stirring at room temperature, followed by (3*R*, 3*aS*, 6*aR*) hexahydrofuro[2,3-*b*]furan-2-yl 4-nitrophenyl carbonate (0.002 g, 0.007 mmol) in one portion as a solid. The yellow reaction mixture was stirred for 9 hours at room temperature. The solvent was removed in vacuo, and the resultant yellow oil was purified by flash column chromatography (1:1 ethyl acetate/hexane) to give *bis*-THF methylenedioxybenzenesulfonamide 1-methyl-3,5-

10 dimethylpyrazole tethered product (56) (0.001 g, 31%) as a colourless oil: R_f = 0.50 (ethyl acetate); coupled LCMS showed the product as a single major peak with m/z 701 $[M+H]^+$; integrated LCMS (204.5 nm) showed the material as a single major peak at R_T of 1.52 min. (purity = 91%).

15

Example 38



20 ***Bis*-THF methylenedioxybenzenesulfonamide ethylpyrazole tethered product (57)**

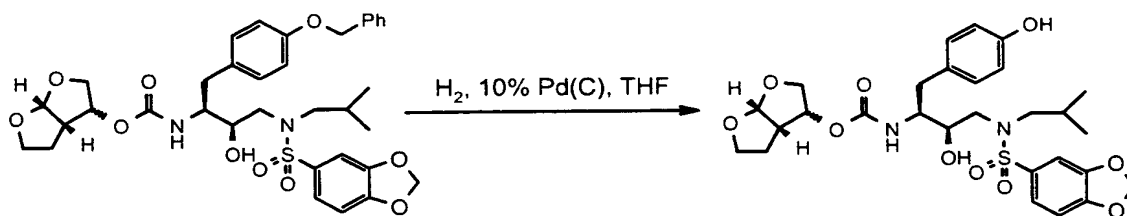
Boc methylenedioxybenzenesulfonamide ethylpyrazole tethered product (0.003 g, 0.005 mmol) was dissolved in DCM (0.15 mL). Trifluoroacetic acid (0.05 mL) was added dropwise, and the reaction was stirred at room temperature for 45 minutes. The solvent was removed in vacuo, and the resultant orange oil was dissolved in DCM

25

(1 mL). Diisopropylethylamine (0.030 mL, 0.18 mmol) was added dropwise with stirring at room temperature, followed by (3R, 3aS, 6aR) hexahydrofuro[2,3-b]furan-2-yl 4-nitrophenyl carbonate (0.002 g, 0.007 mmol) in one
5 portion as a solid. The yellow reaction mixture was stirred for 9 hours at room temperature. The solvent was removed *in vacuo*, and the resultant yellow oil was purified by flash column chromatography (1:1 ethyl acetate/hexane) to give *bis*-THF
10 methylenedioxybenzenesulfonamide ethylpyrazole tethered product (57) (0.001 g, 31%) as a colourless oil: R_f = 0.30 (ethyl acetate); coupled LCMS showed the product as a single major peak with m/z 687 $[M+H]^+$; integrated LCMS (204.5 nm) showed the material as a single major peak at
15 R_T of 1.49 min. (purity = 94%).

006090"7976500

Example 39



(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-

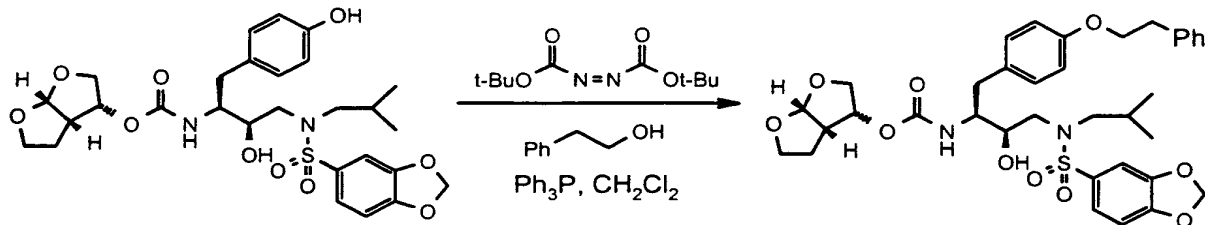
5 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-hydroxybenzyl)propylcarbamate (201)

A solution of 8.00 g (11.7 mmol) of (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-

10 (benzyloxy)benzyl]-2-hydroxypropylcarbamate in 200 mL of THF was subjected to hydrogenation at 50 psi in the presence of 8.0 g of 10% palladium on carbon (Degussa type). After 18 hours the reaction vessel was purged with nitrogen, catalyst removed by filtration through celite, and the filtrate concentrated at reduced pressure to a volume of 30 mL. The solution was rapidly stirred with addition of 30 mL of EtOAc followed by 250 mL of hexane. A white suspension resulted which was stirred at

15 filtration and dried *in vacuo* to afford 6.83 g (98%) of the desired compound as a white powder. ¹H NMR (DMSO-*d*₆): 9.00 (s, 1H), 7.25 (d, 1H), 7.17 (s, 1H), 7.12 (d, 1H), 7.01 (d, 1H), 6.92 (d, 2H), 6.52 (d, 2H), 6.12 (s, 2H), 5.46 (d, 1H), 4.93 (d, 1H), 4.80 (q, 1H), 3.80 (dd, 1H), 20 3.70 (t, 1H), 3.52 (m, 3H), 3.40 (m, 1H), 3.25 (m, 1H), 3.01-2.62 (m, 5H), 2.28 (t, 1H), 1.89 (m, 1H), 1.38 (m, 1H), 1.22 (m, 1H), 0.80 (d, 3H), 0.72 (d, 3H). MS(ESI): 593 (M+H).

Example 40



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-

5 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-

hydroxy-1-[4-(phenethyloxy)benzyl]propylcarbamate (202)

To a solution of 66 mg (0.25 mmol) of triphenylphosphine and 30 μ L (0.25 mmol) of phenethyl alcohol in 3 mL of anhydrous CH_2Cl_2 was added 58 mg (0.25 mmol) of di-tert-

10 butyl azodicarboxylate. The resulting solution was stirred at RT for 5 minutes and was then treated with a solution of 50 mg (0.084 mmol) of

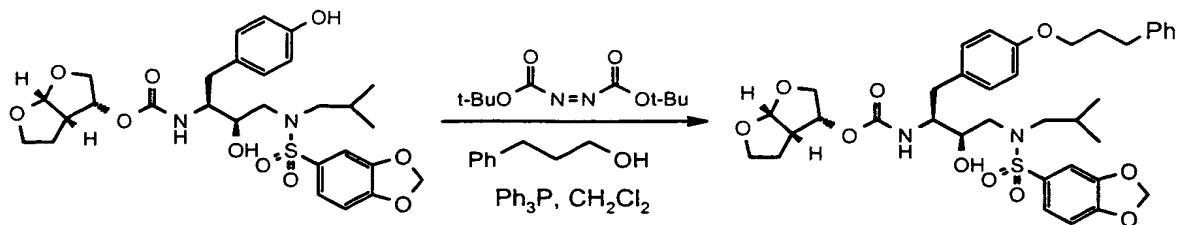
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-

15 hydroxy-1-(4-hydroxybenzyl)propylcarbamate in 2 mL of CH_2Cl_2 . After stirring at RT for 1.5 hours the solution was concentrated to dryness and the residue purified by flash chromatography (SiO_2 , 4:6 hexane/EtOAc) to give the desired product as a white foam in 72% yield. ^1H NMR

20 (CDCl_3): 7.34-7.17 (m, 6H), 7.15 (s, 1H), 7.07 (d, 2H), 6.86 (d, 1H), 6.78 (d, 2H), 6.03 (s, 2H), 5.61 (d, 1H), 4.98 (m, 1H), 4.86 (d, 1H), 4.09 (t, 2H), 3.92 (m, 1H), 3.84-3.56 (m, 6H), 3.17-3.01 (m, 3H), 3.00-2.81 (m, 4H), 2.80-2.64 (m, 2H), 1.78 (m, 1H), 1.56 (m, 1H), 1.47 (m,

25 1H), 0.91 (d, 3H), 0.85 (d, 3H). MS(ESI): 697(M+H).

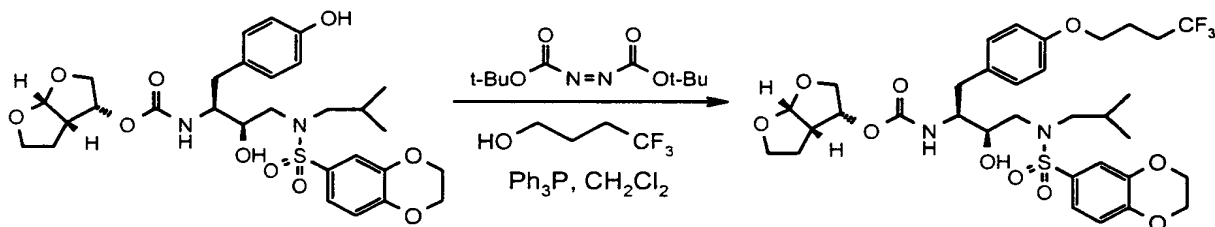
Example 41



(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 5 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-([3-phenylpropyl]oxy)benzyl]propylcarbamate
 (203)

The title compound was prepared according to example 202
 with the exception that 3-phenyl-1-propanol was used
 10 instead of phenethyl alcohol. ¹H NMR (CDCl₃): 7.33-7.11
 (m, 7H), 7.07 (d, 2H), 6.86 (d, 1H), 6.76 (d, 2H), 6.04
 (s, 2H), 5.61 (d, 1H), 4.99 (q, 1H), 4.86 (d, 1H), 3.97-
 3.72 (m, 7H), 3.65 (m, 2H), 3.09 (dd, 1H), 3.01-2.81 (m,
 4H), 2.80-2.64 (m, 4H), 2.06 (m, 2H), 1.85-1.40 (m, 3H),
 15 0.91 (d, 3H), 0.84 (d, 3H). MS(ESI): 711(M+H).

Example 42

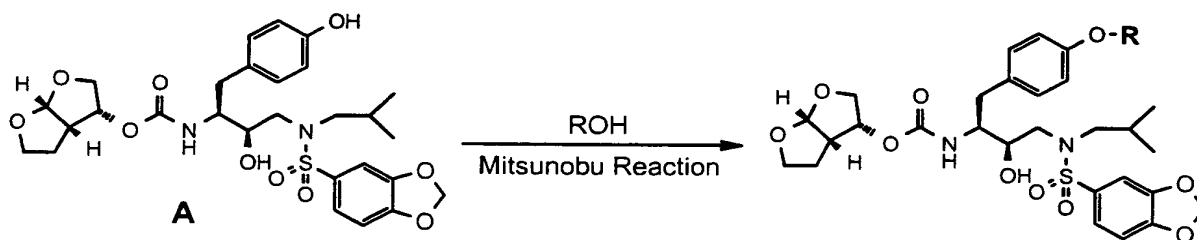


(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 20 [(2,3-dihydro-1,4-benzodioxin-6-
 ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-[4-(4,4,4-
 trifluorobutoxy)benzyl]propylcarbamate (204)

The title compound was prepared according to example 202
 25 with the exception that 4,4,4-trifluorobutanol was used
 instead of phenethyl alcohol. ¹H NMR (CDCl₃): 7.28-7.16

(m, 2H), 7.08 (d, 2H), 6.91 (d, 1H), 6.75 (d, 2H), 5.61 (d, 1H), 5.04-4.85 (m, 2H), 4.25 (m, 4H), 3.91 (m, 3H), 3.88-3.51 (m, 9H), 3.07 (m, 1H), 3.10-2.82 (m, 4H), 2.81-2.65 (m, 2H), 2.26 (m, 2H), 2.00 (m, 2H), 1.84-1.43 (m, 3H), 0.90 (d, 3H), 0.82 (d, 3H). MS(ESI): 717(M+H).

Example 43



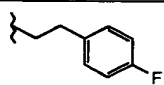
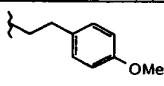
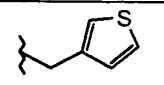
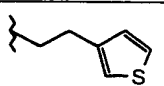
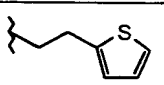
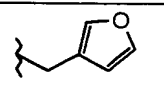
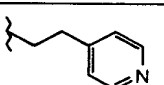
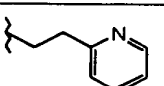
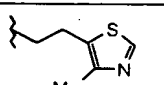
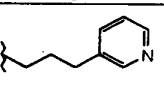
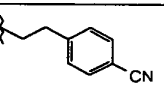
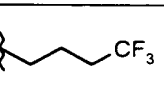
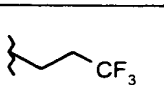
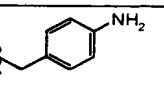
General Procedure for Mitsunobu Alkylations of phenol scaffold A with various alcohols (205-218):

A solution of 2-5 equivalents each of triphenylphosphine and the appropriate alcohol in anhydrous dichloromethane (typically at a concentration of 0.05 M) was treated with di-*t*-butyl azodicarboxylate (2-5 equivalents). Polymer supported triphenylphosphine (Aldrich Chemical) was employed for examples 15-18 to facilitate removal of the triphenylphosphine oxide by-product. After stirring at RT for 5 minutes the solution was treated with 1 equivalent of solid (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-hydroxybenzyl)propylcarbamate and stirring at RT was continued. When the reaction was determined to be complete by thin layer chromatography (2-18 hours) the solution was concentrated in vacuo and the residue purified by flash chromatography (silica gel,

hexane/EtOAc or dichloromethane/2M NH₃ in MeOH) to afford the desired product.

Mass Spectral Data for Compounds 205-218:

5

Example	R	MS (ESI)
205		715 (M+H)
206		727 (M+H)
207		689 (M+H)
208		703 (M+H)
209		703 (M+H)
210		673 (M+H)
211		698 (M+H)
212		698 (M+H)
213		718 (M+H)
214		712 (M+H)
215		722 (M+H)
216		703 (M+H)
217		689 (M+H)
218		698 (M+H)

Proton NMR data for selected compounds from the above table:

5

Example (Compound 209) ^1H NMR(CDCl_3): 7.28 (d, 1H), 7.17-7.04 (m, 4H), 6.95-6.76 (m, 5H), 6.03 (s, 2H), 5.61 (d, 1H), 5.00 (q, 1H), 4.85 (d, 1H), 4.10 (t, 2H), 3.92 (dd, 1H), 3.79 (m, 3H), 3.71-3.48 (m, 3H), 3.24 (t, 2H), 3.10 (m, 1H), 3.01-2.82 (m, 4H), 2.72 (m, 2H), 1.78 (m, 1H), 1.65-1.42 (m, 2H), 0.90 (d, 3H), 0.84 (d, 3H).

Example (Compound 211) ^1H NMR(CDCl_3): 8.53 (br s, 2H), 7.28 (m, 3H), 7.13 (d, 1H), 7.09 (d, 2H), 6.84 (d, 1H), 6.76 (d, 2H), 6.03 (s, 2H), 5.61 (d, 1H), 5.01-4.85 (m, 2H), 4.15 (t, 2H), 3.92 (dd, 1H), 3.85-3.55 (m, 6H), 3.08 (m, 3H), 3.01-2.81 (m, 4H), 2.72 (m, 2H), 1.79 (m, 1H), 1.64-1.41 (m, 2H), 0.89 (d, 3H), 0.82 (d, 3H).

Example (Compound 213) ^1H NMR(CDCl_3): 8.60 (s, 1H), 7.29 (d, 1H), 7.12 (s, 1H), 7.09 (s, 2H), 6.83 (d, 1H), 6.78 (d, 2H), 6.03 (s, 2H), 5.61 (d, 1H), 4.99 (q, 1H), 4.89 (d, 1H), 4.05 (t, 2H), 3.92 (dd, 1H), 3.78 (m, 3H), 3.63 (m, 3H), 3.20 (t, 2H), 3.08 (m, 1H), 3.01-2.81 (m, 4H), 2.72 (m, 2H), 2.41 (s, 3H), 1.78 (m, 1H), 1.58 (m, 1H), 1.46 (m, 1H), 0.89 (s, 3H), 0.82 (s, 3H).

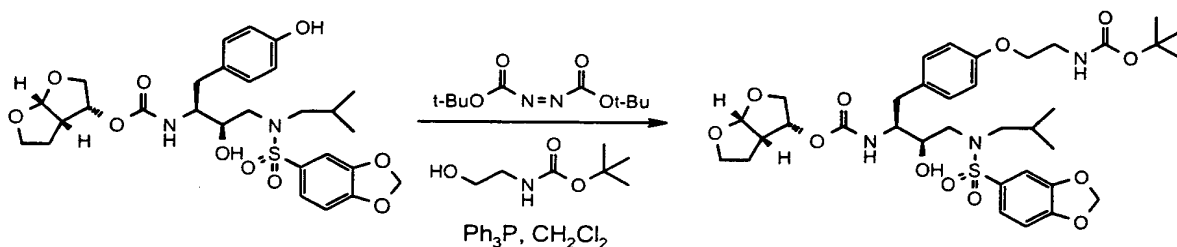
Example (Compound 214) ^1H NMR(CDCl_3): 8.42 (br s, 2H), 7.56 (d, 1H), 7.34-7.19 (m, 2H), 7.13 (s, 1H), 7.07 (d, 2H), 6.85 (d, 1H), 6.73 (d, 2H), 6.03 (s, 2H), 5.60 (d, 1H), 5.10-4.89 (m, 2H), 3.98-3.58 (m, 9H), 3.14-2.65 (m, 9H), 2.08 (m, 2H), 1.80 (m, 1H), 1.61 (m, 1H), 1.50 (m, 1H), 0.89 (d, 3H), 0.81 (d, 3H).

Example (Compound 215) ^1H NMR(CDCl_3): 7.56 (d, 2H), 7.35 (d, 2H), 7.27 (d, 1H), 7.14 (s, 1H), 7.07 (d, 2H), 6.83 (d, 1H), 6.73 (d, 2H), 6.02 (s, 2H), 5.61 (d, 1H), 4.98 (q, 1H), 4.89 (d, 1H), 4.12 (t, 2H), 3.97-3.58 (m, 7H), 3.08 (m, 3H), 3.01-2.81 (m, 4H), 2.81-2.63 (m, 2H), 1.79 (m, 1H), 1.62-1.38 (m, 2H), 0.89 (d, 3H), 0.83 (d, 3H).

Example (Compound 216) ^1H NMR(CDCl_3): 7.38 (d, 1H), 7.21 (s, 1H), 7.17 (d, 2H), 6.94 (d, 1H), 6.84 (d, 2H), 6.12 (s, 2H), 5.69 (d, 1H), 5.13-4.92 (m, 2H), 4.01 (m, 3H), 3.88 (m, 3H), 3.74 (m, 3H), 3.16 (m, 1H), 3.10-2.89 (m, 4H), 2.81 (m, 2H), 2.43-2.20 (m, 2H), 2.08 (m, 2H), 1.94-1.50 (m, 3H), 0.98 (d, 3H), 0.92 (d, 3H).

Example (Compound 217) ^1H NMR(CDCl_3): 7.29 (d, 1H), 7.12 (m, 3H), 6.84 (d, 1H), 6.79 (d, 2H), 6.04 (s, 2H), 5.62 (d, 1H), 5.00 (q, 1H), 4.89 (d, 1H), 4.12 (t, 2H), 3.92 (dd, 1H), 3.80 (m, 2H), 3.66 (m, 2H), 3.12 (dd, 1H), 3.01-2.83 (m, 4H), 2.75 (m, 2H), 2.56 (m, 2H), 1.83-1.44 (m, 5H), 0.91 (d, 3H), 0.83 (d, 3H).

Example 44

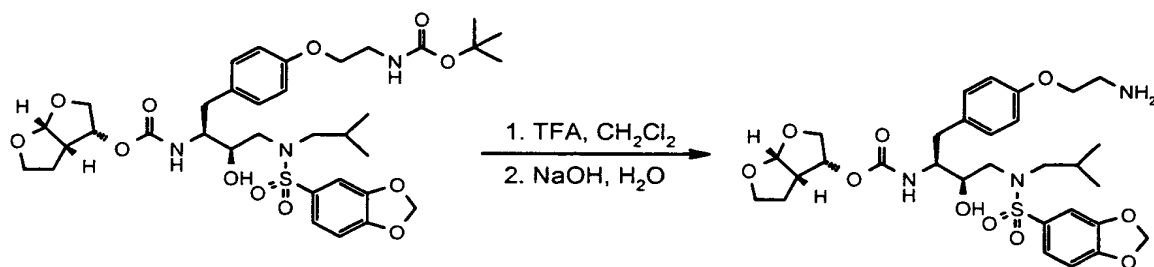


(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-(4-{2-[(*tert*-butoxycarbonyl)amino]ethoxy}benzyl)-2-hydroxypropylcarbamate (219)

The title compound was prepared according to example 202 with the exception that *tert*-butyl 2-hydroxyethylcarbamate was used instead of phenethyl alcohol. ¹H NMR (CDCl₃): 7.30 (dd, 1H), 7.10 (m, 3H), 6.86 (d, 1H), 6.77 (d, 2H), 6.04 (s, 2H), 5.61 (d, 1H), 5.03-4.82 (m, 3H), 3.92 (m, 3H), 3.80 (m, 3H), 3.68 (m, 2H), 3.57 (s, 1H), 3.48 (m, 2H), 3.08 (dd, 1H), 2.92 (m, 4H), 2.75 (m, 2H), 1.78 (m, 1H), 1.64 (m, 1H), 1.41 (m, 10H), 0.90 (d, 3H), 0.83 (d, 3H). MS (ESI): 736 (M+H).

10

Example 45

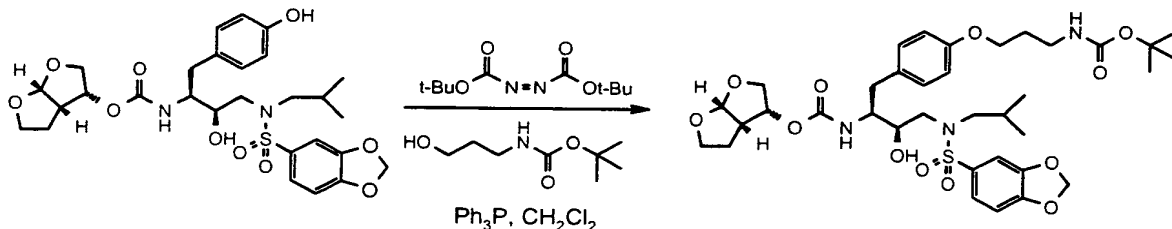


(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-(2-aminoethoxy)benzyl]-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate (220)

A solution of 0.65 g (0.89 mmol) of (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-(4-{2-[(*tert*-butoxycarbonyl)amino]ethoxy}benzyl)-2-hydroxypropylcarbamate in 10 mL of anhydrous CH₂Cl₂ was treated with 15 mL of TFA and the resulting solution was stirred at RT. After 2 hours the solution was evaporated at reduced pressure and the residue redissolved in CH₂Cl₂. The solution was washed with 0.5 M aqueous NaOH (1x), water (2x), dried over MgSO₄, and concentrated *in vacuo*. The crude product was purified by flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 0.40 g (70%)

of the desired compound as a white foam. ¹H NMR (CDCl₃): 7.39 (dd, 1H), 7.12 (s, 1H), 7.08 (d, 2H), 6.84 (d, 1H), 6.77 (d, 2H), 6.03 (s, 2H), 5.61 (d, 1H), 4.95 (m, 2H), 4.00-3.60 (m, 8H), 3.16-2.62 (m, 10H), 2.20-1.40 (m, 5H), 0.90 (d, 3H), 0.82 (d, 3H). MS(ESI): 636(M+H).

Example 46

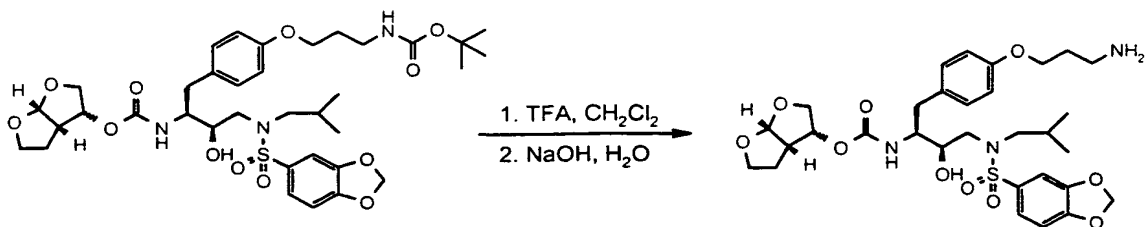


(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-(4-{3-[(tert-butoxycarbonyl)amino]propoxy}benzyl)-2-hydroxypropylcarbamate (221)

The title compound was prepared according to example 202 with the exception that tert-butyl 3-

hydroxypropylcarbamate was used instead of phenethyl alcohol. ¹H NMR (CDCl₃): 7.30 (d, 1H), 7.13 (s, 1H), 7.08 (d, 2H), 6.84 (d, 1H), 6.76 (d, 2H), 6.06 (s, 2H), 5.61 (d, 1H), 5.00 (q, 1H), 4.88 (d, 1H), 4.70 (br s, 1H), 3.93 (m, 3H), 3.80 (m, 4H), 3.67 (m, 2H), 3.26 (m, 2H), 3.09 (dd, 1H), 2.92 (m, 4H), 2.72 (m, 2H), 1.91 (m, 2H), 1.79 (m, 1H), 1.62 (m, 1H), 1.51 (m, 1H), 1.40 (s, 9H), 0.90 (d, 3H), 0.83 (d, 3H). MS(ESI): 750(M+H).

Example 47



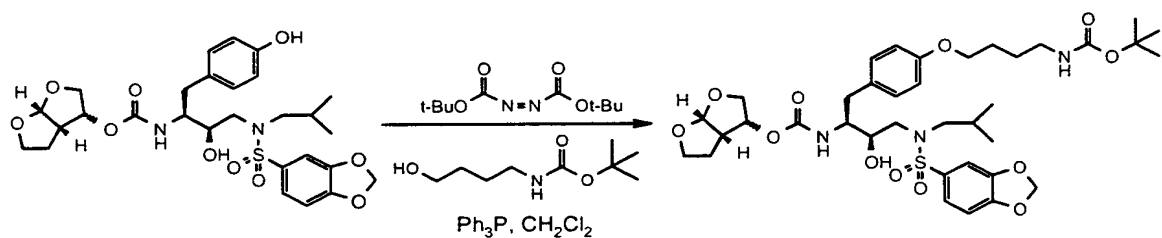
(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-(3-aminopropoxy)benzyl]-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate
(222)

The title compound was prepared according to example 202.

¹H NMR (CDCl₃): 7.29 (d, 1H), 7.15 (s, 1H), 7.09 (d, 2H), 6.85 (d, 1H), 6.78 (d, 2H), 6.04 (s, 2H), 5.61 (d, 1H), 5.00 (m, 2H), 4.03-3.87 (m, 4H), 3.80 (m, 3H), 3.66 (m, 2H), 3.14-2.40 (m, 11H), 1.94 (m, 2H), 1.80 (m, 1H), 1.62 (m, 1H), 1.49 (m, 1H), 0.90 (d, 3H), 0.82 (d, 3H).

MS (ESI): 650 (M+H).

Example 48



(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-(4-{4-[(tert-butoxycarbonyl)amino]butoxy}benzyl)-2-hydroxypropylcarbamate (223)

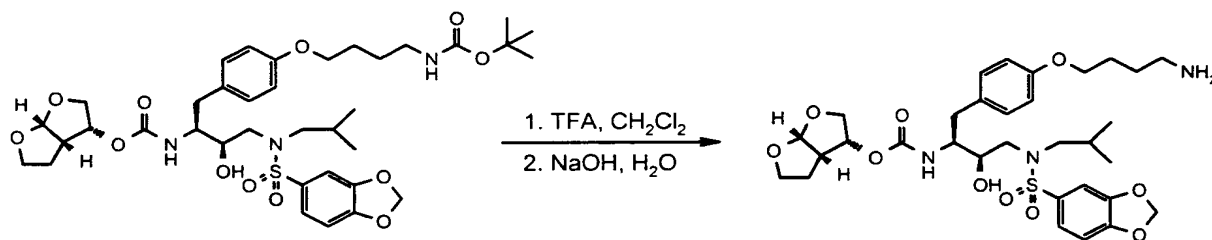
The title compound was prepared according to example 202

with the exception that tert-butyl 3-

hydroxybutylcarbamate (prepared by reaction of 4-amino-1-

butanol with di-*tert*-butyl-dicarbonate in CH_2Cl_2) was used instead of phenethyl alcohol. ^1H NMR (CDCl_3): 7.37 (d, 1H), 7.21 (s, 1H), 7.14 (d, 2H), 6.93 (d, 1H), 6.84 (d, 2H), 6.13 (s, 2H), 5.69 (d, 1H), 5.08 (q, 1H), 4.95 (d, 1H), 4.63 (br s, 1H), 4.08-3.63 (m, 9H), 3.30-2.72 (m, 9H), 1.96-1.40 (m, 16H), 0.97 (d, 3H), 0.90 (d, 3H). MS (ESI): 764 (M+H).

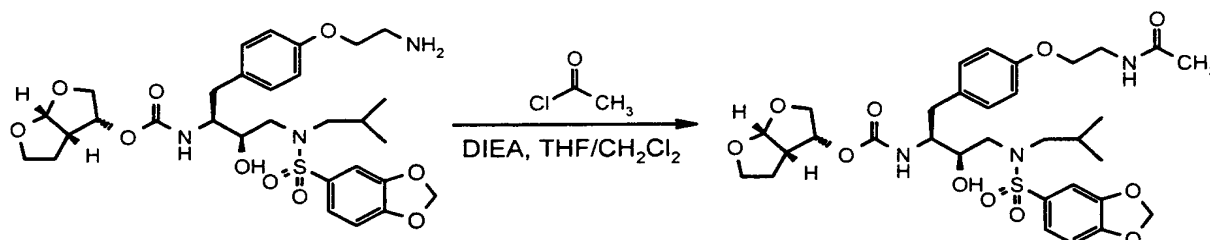
Example 49



(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-(4-aminobutoxy)benzyl]-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate (224)

The title compound was prepared according to example 202. ^1H NMR (CDCl_3): 7.30 (d, 1H), 7.13 (s, 1H), 7.06 (d, 2H), 6.85 (d, 1H), 6.75 (d, 2H), 6.04 (s, 2H), 5.60 (d, 1H), 5.00 (m, 2H), 3.97-3.71 (m, 7H), 3.66 (m, 2H), 3.22-2.60 (m, 11H), 1.87-1.53 (m, 6H), 1.43 (m, 1H), 0.88 (d, 3H), 0.82 (d, 3H). MS (ESI): 664 (M+H).

Example 50

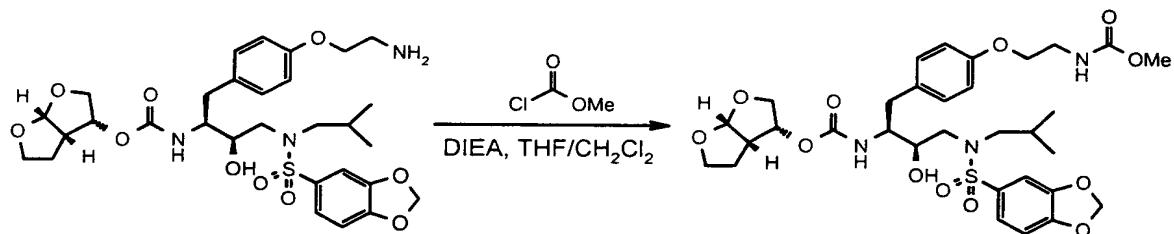


(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-

[2-(acetylamino)ethoxy]benzyl}-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate (225)

A solution of 21 mg (0.033 mmol) of (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-(2-aminoethoxy)benzyl]-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate in 2 mL of 1:1 THF/CH₂Cl₂ was treated with 9 μL (0.050 mmol) of *N,N*-diisopropylethylamine followed by 2.6 μL (0.036 mmol) of acetyl chloride. The resulting solution was stirred at RT. After 1.5 hours the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/MeOH) to afford the desired compound in 86 % yield as a white foam. ¹H NMR (CDCl₃): 7.29 (dd, 1H), 7.16-7.04 (m, 3H), 7.05 (d, 1H), 6.76 (d, 2H), 6.05 (s, 2H), 5.91 (br s, 1H), 5.01 (d, 1H), 5.05-4.89 (m, 2H), 3.94 (m, 3H), 3.80 (m, 3H), 3.72-3.51 (m, 5H), 3.08 (dd, 1H), 3.01-2.83 (m, 4H), 2.74 (m, 2H), 1.97 (s, 3H), 1.86-1.45 (m, 3H), 0.90 (d, 3H), 0.84 (d, 3H). MS(ESI): 678 (M+H).

Example 51



(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-

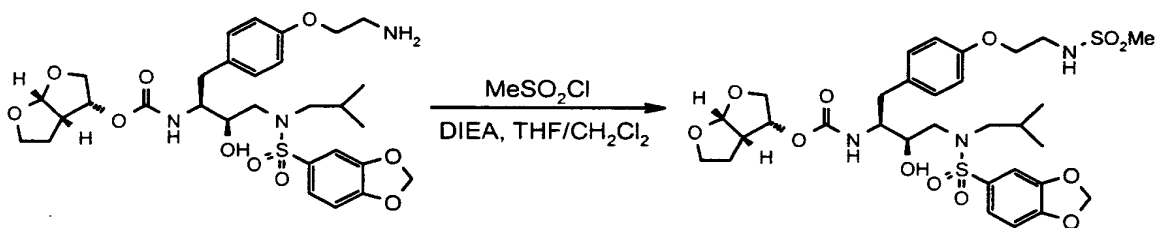
hydroxy-1-(4-{2-[(methoxycarbonyl)amino]ethoxy}
benzyl)propylcarbamate (226)

The title compound was prepared according to example 225
with the exception that methyl chloroformate was used

5 instead of acetyl chloride. ¹H NMR (CDCl₃): 7.29 (dd, 1H),
7.16-7.02 (m, 3H), 6.84 (d, 1H), 6.76 (d, 2H), 6.04 (s,
2H), 5.61 (d, 1H), 5.18-4.84 (m, 3H), 4.02-3.43 (m, 14H),
3.09 (m, 1H), 2.91 (m, 4H), 2.72 (m, 2H), 1.85-1.43 (m,
3H), 0.89 (d, 3H), 0.92 (d, 3H). MS(ESI): 694 (M+H).

10

Example 52



15

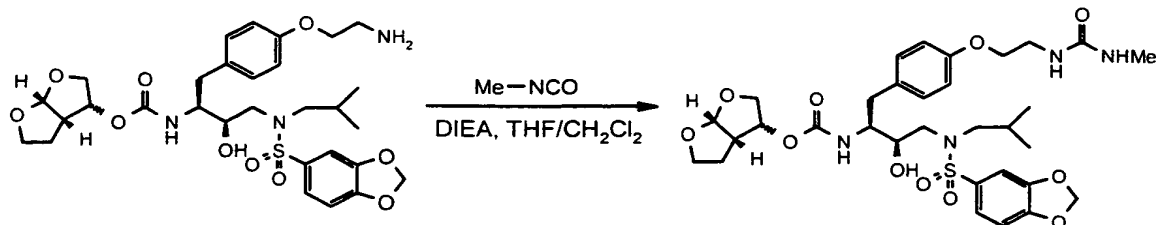
(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-(4-{2-[(methylsulfonyl)amino]ethoxy}

20 benzyl)propylcarbamate (227)

The title compound was prepared according to example 225
with the exception that methanesulfonyl chloride was used
instead of acetyl chloride. ¹H NMR (CDCl₃): 7.29 (dd, 1H),

25 7.11 (m, 3H), 6.86 (d, 1H), 6.77 (d, 2H), 6.04 (s, 2H),
5.61 (d, 1H), 4.98 (m, 2H), 4.84 (m, 1H), 4.09-3.44 (m,
11H), 3.12-2.82 (m, 8H), 2.74 (m, 2H), 1.88-1.43 (m, 3H),
0.89 (d, 3H), 0.81 (d, 3H). MS(ESI): 714 (M+H).

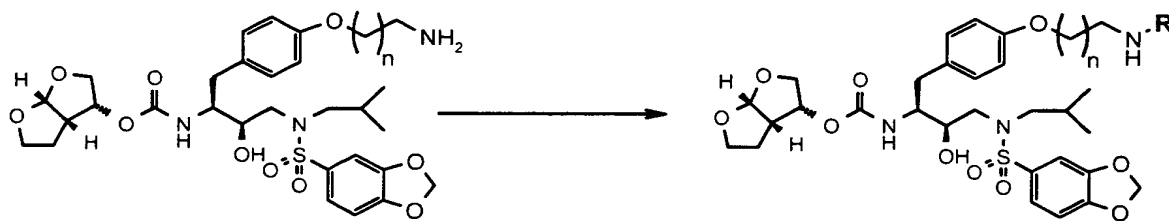
Example 53



5 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-(2-{[(methylamino)carbonyl]amino}
 ethoxy)benzyl]propylcarbamate (228)

The title compound was prepared according to example 225
 10 with the exception that methyl isocyanate was used
 instead of acetyl chloride. ¹H NMR (CDCl₃): 7.29 (dd, 1H),
 7.13 (s, 1H), 7.08 (d, 2H), 6.84 (d, 1H), 6.76 (d, 2H),
 6.05 (s, 2H), 5.61 (d, 1H), 5.10-4.90 (m, 2H), 4.04-3.48
 (m, 11H), 3.15-2.67 (m, 10H), 1.80 (m, 1H), 1.62 (m, 1H),
 15 1.47 (m, 1H), 0.85 (m, 6H). MS(ESI): 693(M+H).

Example 54

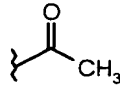
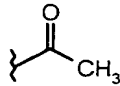
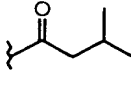
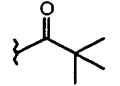
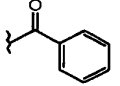
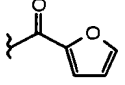
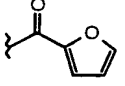
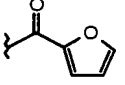
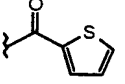


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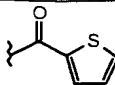
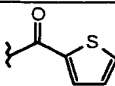
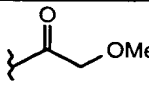
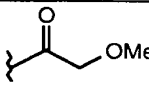
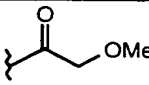
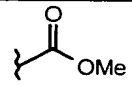
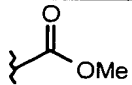
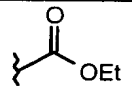
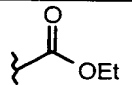
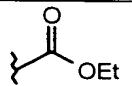
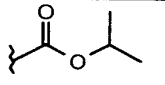
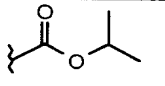
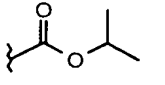
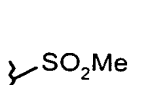
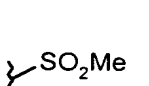
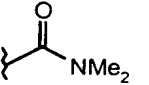
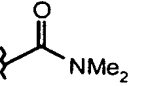
General procedure for reactions of primary amine
 scaffolds with various electrophiles (229-260): A
 solution of the primary amine (0.02 M) in anhydrous CH₂Cl₂
 at 0°C was treated with 1.5 equivalents of N,N-
 25 diisopropylethylamine (omitted for reactions with

isocyanates and isothiocyanates, examples 56-60) followed by 1.05 equivalent of the appropriate electrophile (acid chloride, chloroformate, sulfonyl chloride, carbamyl chloride, isocyanate, isothiocyanate). The resulting solution was allowed to warm to RT with stirring. When analysis by TLC indicated the reaction to be complete (2-18 hours) the solution was concentrated *in vacuo* and the residue subjected to flash chromatography (SiO₂, CH₂Cl₂/MeOH or hexane/EtOAc) to afford the desired product.

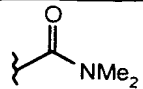
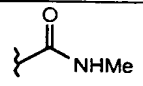
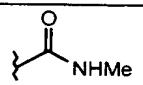
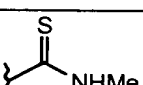
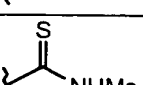
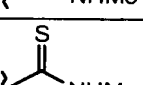
Mass Spectral Data for Compounds 29-60:

Example	n	R	MS (ESI)
229	2		692 (M+H)
230	3		706 (M+H)
231	2		734 (M+H)
232	2		734 (M+H)
233	2		754 (M+H)
234	1		730 (M+H)
235	2		744 (M+H)
236	3		758 (M+H)
237	1		746 (M+H)

000000-1947650

238	2		760 (M+H)
239	3		774 (M+H)
240	1		708 (M+H)
241	2		722 (M+H)
242	3		736 (M+H)
243	2		708 (M+H)
244	3		722 (M+H)
245	1		708 (M+H)
246	2		722 (M+H)
247	3		736 (M+H)
248	1		722 (M+H)
249	2		736 (M+H)
250	3		750 (M+H)
251	2		728 (M+H)
252	3		742 (M+H)
253	1		707 (M+H)
254	2		721 (M+H)

000000-131650

255	3		735 (M+H)
256	2		707 (M+H)
257	3		721 (M+H)
258	1		709 (M+H)
259	2		723 (M+H)
260	3		737 (M+H)

Proton NMR data for selected compounds from the above table:

5

Example (Compound 231) NMR(CDCl₃): 7.29 (d, 1H), 7.13 (s, 1H), 7.08 (d, 2H), 6.86 (d, 1H), 6.75 (d, 2H), 6.04 (s, 2H), 5.80 (br s, 1H), 5.60 (d, 1H), 4.98 (m, 2H), 3.92 (m, 3H), 3.79 (m, 3H), 3.65 (m, 2H), 3.40 (q, 2H), 3.15-2.65 (m, 8H), 2.13-1.90 (m, 5H), 1.80 (m, 1H), 1.62 (m, 1H), 1.49 (m, 1H), 0.95-0.78 (m, 12H).

10

Example (Compound 233) NMR(CDCl₃): 7.72 (d, 2H), 7.43 (d, 1H), 7.38 (m, 2H), 7.29 (d, 1H), 7.10 (m, 3H), 6.84 (d, 1H), 6.77 (d, 2H), 6.65 (br s, 1H), 6.03 (s, 2H), 5.59 (d, 1H), 4.98 (m, 2H), 4.02 (t, 2H), 3.91 (dd, 1H), 3.79 (m, 3H), 3.63 (m, 5H), 3.14-2.66 (m, 7H), 2.07 (m, 2H), 1.79 (m, 1H), 1.61 (m, 1H), 1.49 (m, 1H), 0.89 (d, 3H), 0.92 (d, 3H).

20

Example (Compound 235) NMR(CDCl₃): 7.41 (s, 1H), 7.28 (d, 1H), 7.18-7.04 (m, 4H), 6.87 (d, 1H), 6.79 (m, 3H), 6.45

(m, 1H), 6.04 (s, 2H), 5.60 (d, 1H), 4.96 (m, 2H), 4.02
(t, 2H), 3.91 (dd, 1H), 3.80 (m, 3H), 3.73-3.52 (m, 5H),
3.08 (m, 1H), 3.01-2.82 (m, 4H), 2.74 (m, 2H), 2.05 (m,
2H), 1.80 (m, 1H), 1.63 (m, 1H), 1.50 (m, 1H), 0.90 (d,
5 3H), 0.82 (d, 3H).

Example (Compound 236) NMR(CDCl₃): 7.38 (s, 1H), 7.29 (d,
1H), 7.13 (s, 1H), 7.07 (m, 3H), 6.84 (d, 1H), 6.77 (d,
2H), 6.46 (m, 2H), 6.03 (s, 2H), 5.60 (d, 1H), 4.96 (m,
10 2H), 3.92 (m, 3H), 3.80 (m, 3H), 3.66 (m, 3H), 3.47 (q,
2H), 3.08 (dd, 1H), 2.92 (m, 4H), 2.73 (m, 2H), 1.80 (m,
5H), 1.62 (m, 1H), 1.48 (m, 1H), 0.90 (d, 3H), 0.82 (d,
3H).

Example (Compound 237) NMR(CDCl₃): 7.50 (m, 1H), 7.43 (d,
1H), 7.29 (d, 1H), 7.12 (m, 3H), 7.02 (m, 1H), 6.85 (d,
1H), 6.80 (d, 2H), 6.43 (br s, 1H), 6.04 (s, 2H), 5.60
(d, 1H), 4.96 (m, 2H), 4.05 (t, 2H), 3.91 (dd, 1H), 3.80
(m, 6H), 3.66 (m, 2H), 3.08 (dd, 1H), 3.00-2.81 (m, 4H),
20 2.73 (m, 2H), 1.80 (m, 1H), 1.62 (m, 1H), 1.50 (m, 1H),
0.89 (d, 3H), 0.82 (d, 3H).

Example (Compound 238) NMR(CDCl₃): 7.47 (d, 1H), 7.42 (d,
1H), 7.30 (d, 1H), 7.11 (m, 3H), 7.03 (t, 1H), 6.85 (t,
25 1H), 6.79 (d, 2H), 6.51 (br s, 1H), 6.04 (s, 2H), 5.61
(d, 1H), 4.98 (m, 2H), 4.02 (t, 2H), 3.91 (dd, 1H), 3.80
(m, 3H), 3.70-3.52 (m, 4H), 3.08 (m, 1H), 3.00-2.83 (m,
5H), 2.81-2.67 (m, 2H), 2.07 (m, 2H), 1.80 (m, 1H), 1.62
(m, 1H), 1.50 (m, 1H), 0.89 (d, 3H), 0.81 (d, 3H).

30

Example (Compound 241) NMR(CDCl₃): 7.29 (d, 1H), 7.12 (s,
1H), 7.08 (d, 2H), 6.89 (br s, 1H), 6.85 (d, 1H), 6.77
(d, 2H), 6.04 (s, 2H), 5.61 (d, 1H), 4.99 (q, 1H), 4.91

(d, 1H), 4.01-3.89 (m, 3H), 3.88-3.72 (m, 6H), 3.67 (m, 2H), 3.46 (q, 2H), 3.39 (s, 3H), 3.09 (dd, 1H), 3.01-2.82 (m, 4H), 2.81-2.67 (m, 2H), 1.98 (m, 2H), 1.80 (m, 1H), 1.63 (m, 1H), 1.50 (m, 1H), 0.89 (d, 3H), 0.82 (d, 3H).

5

Example (Compound 243) NMR(CDCl₃): 7.38 (d, 1H), 7.20 (s, 1H), 7.16 (d, 2H), 6.92 (d, 1H), 6.85 (d, 2H), 6.12 (s, 2H), 5.69 (d, 1H), 5.14-4.89 (m, 3H), 4.01 (m, 3H), 3.89 (m, 3H), 3.73 (m, 6H), 3.42 (m, 2H), 3.18 (dd, 1H), 3.11-2.90 (m, 4H), 2.82 (m, 2H), 2.01 (m, 2H), 1.93-1.52 (m, 3H), 0.98 (d, 3H), 0.92 (d, 3H).

Example (Compound 244) NMR(CDCl₃): 7.38 (d, 1H), 7.22 (s, 1H), 7.14 (d, 2H), 6.93 (d, 1H), 6.85 (d, 2H), 6.13 (s, 2H), 5.69 (d, 1H), 5.12-4.90 (m, 2H), 4.82 (br s, 1H), 4.07-3.61 (m, 12H), 3.36-3.10 (m, 3H), 3.09-2.90 (m, 4H), 2.90-2.70 (m, 2H), 1.95-1.62 (m, 6H), 1.56 (m, 1H), 0.98 (d, 3H), 0.91 (d, 3H).

Example (Compound 247) NMR(CDCl₃): 7.29 (d, 1H), 7.13 (s, 1H), 7.07 (d, 2H), 6.84 (d, 1H), 6.76 (d, 2H), 6.06 (s, 2H), 5.61 (d, 1H), 5.02-4.84 (m, 2H), 4.71 (br s, 1H), 4.08 (q, 2H), 3.97-3.73 (m, 7H), 3.65 (m, 2H), 3.20 (m, 2H), 3.08 (dd, 1H), 3.00-2.82 (m, 4H), 2.73 (m, 2H), 1.78 (m, 3H), 1.64 (m, 3H), 1.49 (m, 1H), 1.20 (t, 3H), 0.89 (d, 3H), 0.82 (d, 3H).

Example (Compound 249) NMR(CDCl₃): 7.29 (d, 1H), 7.12 (s, 1H), 7.09 (d, 2H), 6.84 (d, 1H), 6.77 (d, 2H), 6.06 (s, 2H), 5.61 (d, 1H), 5.04-4.70 (m, 4H), 3.94 (m, 3H), 3.80 (m, 3H), 3.67 (m, 2H), 3.32 (m, 2H), 3.07 (dd, 1H), 2.91 (m, 5H), 2.75 (m, 2H), 1.94 (m, 2H), 1.80 (m, 1H), 1.63

006090" 494T6560

(m, 1H), 1.51 (m, 1H), 1.20 (d, 6H), 0.90 (d, 3H), 0.82 (d, 3H).

5 Example (Compound 250) NMR(CDCl₃): 7.29 (d, 1H), 7.12 (s, 1H), 7.02 (d, 2H), 6.84 (d, 1H), 6.76 (d, 2H), 6.03 (s, 2H), 5.61 (d, 1H), 5.00 (q, 1H), 4.88 (m, 2H), 4.65 (br s, 1H), 3.90 (m, 3H), 3.80 (m, 3H), 3.66 (m, 2H), 3.20 (m, 2H), 3.10 (dd, 1H), 2.91 (m, 5H), 2.73 (m, 2H), 1.78 (m, 3H), 1.62 (m, 3H), 1.48 (m, 1H), 1.20 (d, 6H), 0.89 (d, 3H), 0.81 (d, 3H).

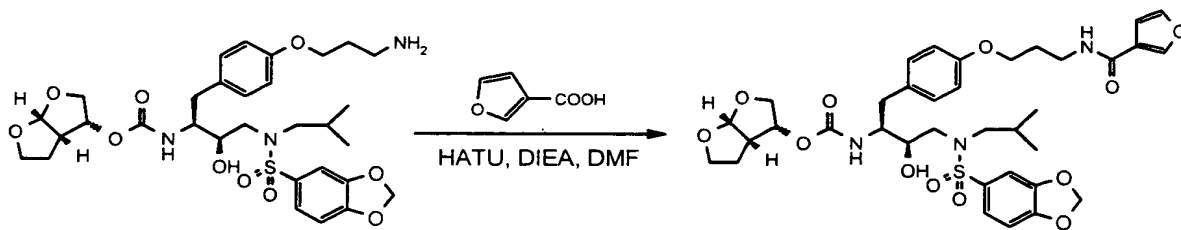
15 Example (Compound 252) NMR(CDCl₃): 7.29 (d, 1H), 7.13 (s, 1H), 7.08 (d, 2H), 6.83 (d, 1H), 6.76 (d, 2H), 6.04 (s, 2H), 5.60 (d, 1H), 4.96 (m, 2H), 4.53 (br s, 1H), 3.92 (m, 3H), 3.79 (m, 3H), 3.67 (m, 2H), 3.16 (m, 2H), 3.07 (dd, 1H), 3.04-2.82 (m, 8H), 2.81-2.65 (m, 2H), 1.87-1.54 (m, 6H), 1.49 (m, 1H), 0.89 (d, 3H), 0.82 (d, 3H).

20 Example (Compound 254) NMR(CDCl₃): 7.29 (d, 1H), 7.12 (s, 1H), 7.08 (d, 2H), 6.85 (d, 1H), 6.74 (d, 2H), 6.04 (s, 2H), 5.60 (d, 1H), 5.00 (m, 2H), 4.82 (br s, 1H), 4.01-3.88 (m, 3H), 3.80 (m, 3H), 3.66 (m, 2H), 3.37 (t, 2H), 3.15-2.82 (m, 12H), 2.81-2.65 (m, 2H), 1.96 (m, 2H), 1.80 (m, 1H), 1.62 (m, 1H), 1.50 (m, 1H), 0.90 (d, 3H), 0.82 (d, 3H).

30 Example (Compound 260) NMR(CDCl₃): 7.29 (d, 1H), 7.13 (s, 1H), 7.07 (d, 2H), 6.86 (d, 1H), 6.75 (d, 2H), 6.04 (s, 2H), 5.60 (d, 1H), 4.98 (m, 2H), 3.90 (m, 3H), 3.80 (m, 3H), 3.65 (m, 2H), 3.48 (br s, 2H), 3.08 (m, 1H), 3.02-2.81 (m, 10H), 2.80-2.62 (m, 2H), 1.80 (m, 5H), 1.61 (m, 1H), 1.48 (m, 1H), 0.88 (d, 3H), 0.81 (d, 3H).

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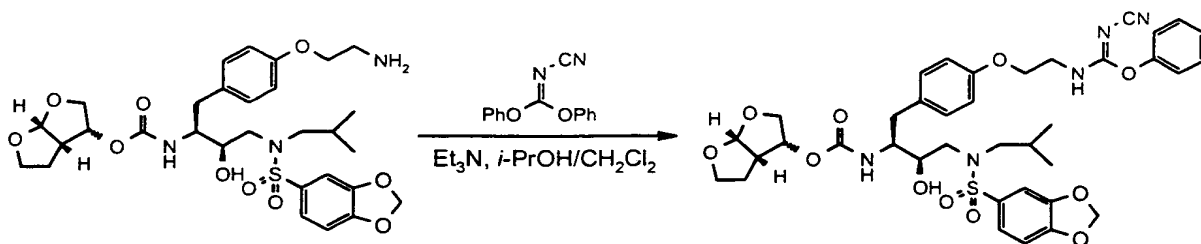
Example 55



5 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[3-
 (3-furoylamino)propoxy]benzyl}-2-hydroxypropylcarbamate
 (261)

A solution of 44 mg (0.068 mmol) of (3R,3aS,6aR)-
 10 hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-1-[4-(3-
 aminopropoxy)benzyl]-3-[(1,3-benzodioxol-5-
 ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate, 15
 mg (0.14 mmol) of 3-furoic acid, and 36 μ L (0.20 mmol) of
 N,N-diisopropylethylamine in 2 mL of anhydrous DMF was
 15 treated with 52 mg (0.14 mmol) of O-(7-azabenzotriazol-1-
 yl)-1,1,3,3-tetramethyluronium hexafluorophosphate (HATU)
 and the resulting solution was stirred at RT. After 18
 hours the solution was concentrated *in vacuo* and the
 residue subjected to flash chromatography (SiO₂, 97:3
 20 CH₂Cl₂/MeOH) to afford 38 mg (75%) of the desired compound
 as a white foam. ¹H NMR (CDCl₃): 7.88 (s, 1H), 7.38 (s,
 1H), 7.29 (d, 1H), 7.10 (m, 3H), 6.84 (d, 1H), 6.76 (d,
 2H), 6.56 (s, 1H), 6.34 (br s, 1H), 6.05 (s, 2H), 5.60
 (d, 1H), 5.00 (m, 2H), 4.01 (t, 2H), 3.90 (dd, 1H), 3.80
 25 (m, 3H), 3.64 (m, 2H), 3.55 (q, 2H), 3.17-2.63 (m, 8H),
 2.03 (m, 2H), 1.80 (m, 1H), 1.61 (m, 1H), 1.49 (m, 1H),
 0.89 (d, 3H), 0.83 (d, 3H). MS(ESI): 744 (M+H).

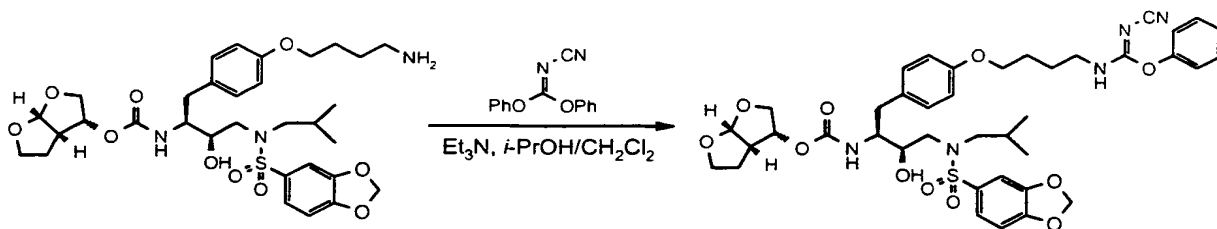
Example 56



(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(2-
 5 {[(*E* and/or *Z*)-(cyanoimino)(phenoxy)methyl]amino}
 ethoxy)benzyl]-2-hydroxypropylcarbamate (262)

A solution of 0.20 g (0.32 mmol) of (3*R*,3*aS*,6*aR*)-
 hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-(2-
 aminoethoxy)benzyl]-3-[(1,3-benzodioxol-5-
 10 ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate, 83
 mg (0.35 mmol) of diphenyl cyanocarbonimidate, and 66 μ L
 of (0.47 mmol) triethylamine in 12 mL of 1:1 *i*-PrOH/ CH_2Cl_2
 was stirred at RT. After 2.5 hours the solution was
 concentrated to dryness at reduced pressure and the
 15 residue subjected to flash chromatography (SiO_2 , 95:5
 $\text{CH}_2\text{Cl}_2/\text{MeOH}$) to afford 0.24 g (96%) of the desired
 compound as a white foam. ^1H NMR (CDCl_3): 7.47-7.21 (m,
 4H), 7.20-7.02 (m, 5H), 6.88-6.63 (m, 3H), 6.42 (br s,
 1H), 6.05 (s, 2H), 5.61 (d, 1H), 4.99 (m, 2H), 4.20-3.50
 20 (m, 11H), 3.09 (m, 1H), 3.01-2.82 (m, 4H), 2.76 (m, 2H),
 1.78 (m, 1H), 1.72-1.43 (m, 2H), 0.90 (d, 3H), 0.81 (d,
 3H). MS(ESI): 780 ($\text{M}+\text{H}$).

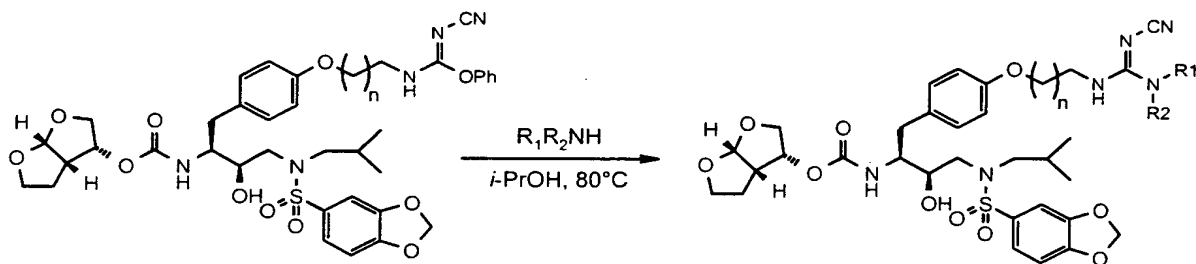
Example 57



(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(4-
 5 {[(E and/or Z)-(cyanoimino)(phenoxy)methyl]amino}
 butoxy)benzyl]-2-hydroxypropylcarbamate (263)

According to example 262, (3R,3aS,6aR)-hexahydrofuro[2,3-
 b]furan-3-yl (1S,2R)-1-[4-(4-aminobutoxy)benzyl]-3-[(1,3-
 benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 10 hydroxypropylcarbamate was converted to the desired
 compound which was obtained in 98% yield as a white foam.
¹H NMR (CDCl₃): 7.52-7.26 (m, 4H), 7.25-7.07 (m, 5H),
 6.97-6.67 (m, 4H), 6.12 (s, 2H), 5.69 (d, 1H), 5.06 (m,
 2H), 4.10-3.40 (m, 11H), 3.22-2.73 (m, 7H), 2.00-1.55 (m,
 15 7H), 0.96 (d, 3H), 0.91 (d, 3H). MS(ESI): 808(M+H).

Example 58

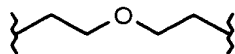


20 General procedure for the synthesis of cyanoguanidines
 from imidocarbamate scaffolds (264-269):

A mixture of 0.05 mmol of imidocarbamate intermediate in
 3 mL of isopropanol in a sealed tube was treated with 10-
 20 equivalents of amine (2M NH₃ in MeOH, 8M MeNH₂ in EtOH,
 25 2M Me₂NH in THF, neat morpholine). Upon heating, the

solid starting material dissolved to give a clear solution. After stirring at 80°C for 3 hours the solution was cooled to RT and concentrated to dryness at reduced pressure. The residue was subjected to flash chromatography (SiO₂, CH₂Cl₂/MeOH or CH₂Cl₂/2M NH₃ in MeOH) to afford the desired compound as white foam.

Mass Spectral Data for Compounds 264-269

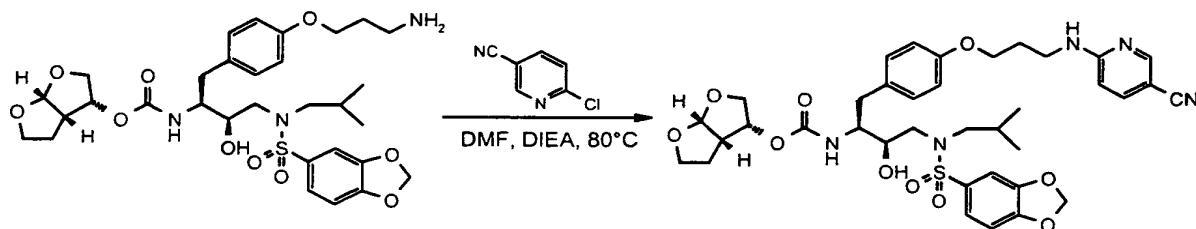
Example	n	R ₁	R ₂	MS (ESI)
264	1	H	H	703 (M+H)
265	1	Me	H	717 (M+H)
266	1			773 (M+H)
267	3	H	H	731 (M+H)
268	3	Me	H	745 (M+H)
269	3	Me	Me	759 (M+H)

Proton NMR data for selected compounds from the above table:

Example (Compound 265) ¹H NMR(CDCl₃): 7.30 (d, 1H), 7.11 (m, 3H), 6.86 (d, 1H), 6.76 (d, 2H), 6.04 (s, 2H), 5.87 (br s, 1H), 5.65-5.42 (m, 2H), 5.09 (d, 1H), 5.00 (q, 1H), 4.01 (m, 2H), 3.92 (dd, 1H), 3.79 (m, 3H), 3.71-3.52 (m, 5H), 3.11-2.66 (10H), 1.80 (m, 1H), 1.61 (m, 1H), 1.50 (m, 1H), 0.86 (m, 6H).

Example (Compound 269) ¹H NMR(CDCl₃): 7.29 (d, 1H), 7.12 (s, 1H), 7.08 (d, 2H), 6.84 (d, 1H), 6.74 (d, 2H), 6.03 (s, 2H), 5.60 (d, 1H), 5.06-4.87 (m, 3H), 3.91 (m, 3H), 3.79 (m, 3H), 3.64 (m, 3H), 3.49 (q, 2H), 3.15-2.65 (m, 13H), 1.89-1.54 (m, 6H), 1.48 (m, 1H), 0.86 (m, 6H).

Example 59



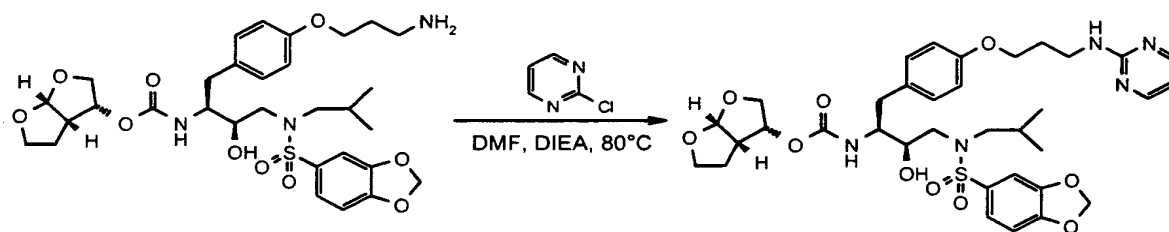
(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-(4-{3-[(5-cyano-2-pyridinyl)amino]propoxy}benzyl)-2-hydroxypropylcarbamate (270)

A solution of 35 mg (0.054 mmol) of (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-1-[4-(3-aminopropoxy)benzyl]-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate, 22 mg (0.16 mmol) of 2-chloro-5-cyanopyridine, and 38 μ L (0.22 mmol) of *N,N*-diisopropylethylamine in 0.5 mL of anhydrous DMF was heated to 80°C with stirring. After 6 hours the solution was cooled to RT and concentrated to dryness *in vacuo*. The residue was purified by flash chromatography (SiO₂, 95:5 CH₂Cl₂/MeOH) to afford 33 mg (80%) of the desired compound as a white foam. ¹H NMR (CDCl₃): 8.30 (s, 1H), 7.50 (d, 1H), 7.29 (d, 1H), 7.13 (s, 1H), 7.09 (d, 2H), 6.84 (d, 1H), 6.78 (d, 2H), 6.37 (d, 1H), 6.04 (s, 2H), 5.62 (d, 1H), 5.44 (br s, 1H), 4.96 (m, 2H), 4.00 (t, 2H), 3.91 (m, 1H), 3.80 (m, 3H), 3.71-3.49 (m, 5H), 3.18-2.65 (m, 7H), 2.08 (m, 2H), 1.80

(m, 1H), 1.61 (m, 1H), 1.50 (m, 1H), 0.90 (d, 3H), 0.81 (d, 3H). MS (ESI): 752 (M+H).

5

Example 60

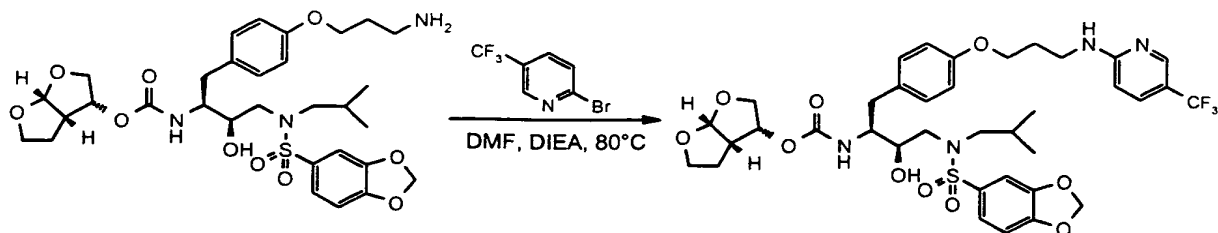


10 (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-{4-[3-(2-pyrimidinylamino)propoxy]benzyl}
propylcarbamate (271)

The title compound was prepared according to example 270
15 with the exception that 2-chloropyrimidine was used
instead of 2-chloro-5-cyanopyridine. ¹H NMR (CDCl₃): 8.23
(m, 2H), 7.29 (d, 1H), 7.12 (s, 1H), 7.06 (d, 2H), 6.86
(d, 1H), 6.80 (d, 2H), 6.50 (t, 1H), 6.06 (s, 2H), 5.60
(d, 1H), 5.50 (br s, 1H), 5.00 (m, 1H), 4.88 (d, 1H),
20 4.00 (t, 2H), 3.92 (m, 1H), 3.80 (m, 3H), 3.71-3.52 (m,
5H), 3.09 (m, 1H), 3.00-2.81 (m, 4H), 2.73 (m, 2H), 2.13-
1.71 (m, 3H), 1.64 (m, 1H), 1.52 (m, 1H), 0.90 (d, 3H),
0.83 (d, 3H). MS (ESI): 728 (M+H).

25

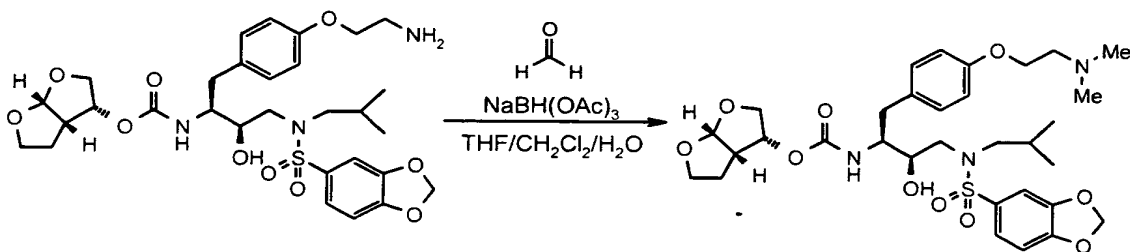
Example 61



(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 5 hydroxy-1-[4-(3-{[5-(trifluoromethyl)-2-
 pyridinyl]amino}propoxy)benzyl]propylcarbamate (272)

The title compound was prepared according to example 270
 with the exception that 2-bromo-5-(trifluoromethyl)
 pyridine was used instead of 2-chloro-5-cyanopyridine. ¹H
 10 NMR (CDCl₃): 8.34 (s, 1H), 7.60 (d, 1H), 7.37 (d, 1H),
 7.21 (s, 1H), 7.16 (d, 2H), 6.93 (d, 1H), 6.86 (d, 2H),
 6.45 (d, 1H), 6.12 (s, 2H), 5.69 (d, 2H), 5.32 (br s,
 1H), 5.07 (m, 2H), 4.15-3.66 (m, 9H), 3.60 (m, 2H), 3.22-
 2.70 (m, 7H), 2.13 (m, 2H), 1.87 (m, 1H), 1.77-1.48 (m,
 15 2H), 0.98 (d, 3H), 0.91 (d, 3H). MS(ESI): 795(M+H).

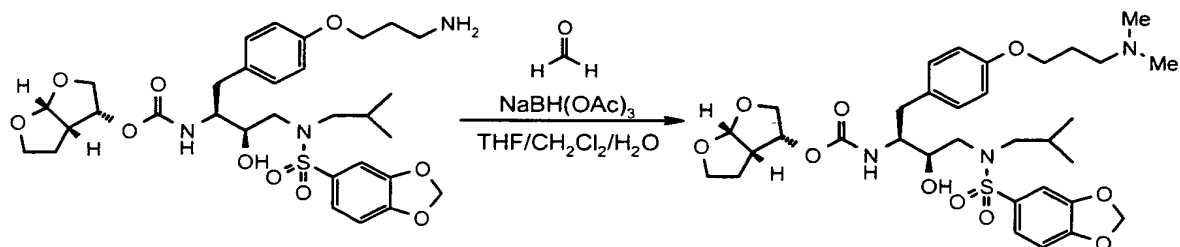
Example 62



(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[2-
 (dimethylamino)ethoxy]benzyl}-2-hydroxypropylcarbamate
 (273)

A solution of 21 mg (0.033 mmol) of (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-1-[4-(2-aminoethoxy)benzyl]-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate in 3 mL of 8:2 THF/CH₂Cl₂ was treated with 13 μL (0.17 mmol) of 37% aqueous formaldehyde followed by 35 mg (0.17 mmol) of NaBH(OAc)₃ and the resulting cloudy solution was stirred at RT. After 3 hours the solution was filtered to remove solids and the filtrate concentrated *in vacuo*. The residue was purified by flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to give the desired compound as a white foam in 68% yield. ¹H NMR (CDCl₃): 7.29 (dd, 1H), 7.13 (s, 1H), 7.07 (d, 2H), 6.85 (d, 1H), 6.80 (d, 2H), 6.03 (s, 2H), 5.61 (d, 1H), 4.98 (q, 1H), 4.89 (d, 1H), 4.01 (t, 2H), 3.92 (m, 1H), 3.80 (m, 3H), 3.66 (m, 2H), 3.09 (dd, 1H), 2.92 (m, 5H), 2.74 (m, 4H), 2.32 (s, 6H), 1.80 (m, 1H), 1.63 (m, 1H), 1.42 (m, 1H), 0.90 (d, 3H), 0.84 (d, 3H). MS(ESI): 664 (M+H).

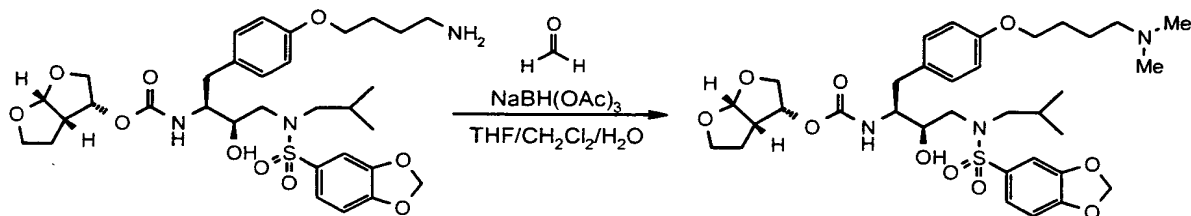
Example 63



(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-[3-(dimethylamino)propoxy]benzyl]-2-hydroxypropylcarbamate (274)

According to example 273, (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-
 b]furan-3-yl (1*S*,2*R*)-1-[4-(3-aminopropoxy)benzyl]-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxypropylcarbamate was subjected to reductive
 5 alkylation with formaldehyde to give the desired compound
 as a white foam in 76% yield. ¹H NMR (CDCl₃): 7.37 (d,
 1H), 7.22 (s, 1H), 7.15 (d, 2H), 6.92 (d, 1H), 6.84 (d,
 2H), 6.13 (s, 2H), 5.70 (d, 1H), 5.07 (q, 1H), 4.95 (d,
 1H), 4.08-3.63 (m, 9H), 3.18 (dd, 1H), 3.00 (m, 4H), 2.82
 10 (m, 2H), 2.50 (t, 2H), 2.31 (s, 6H), 2.10-1.50 (m, 5H),
 0.97 (d, 3H), 0.91 (d, 3H). MS(ESI): 678 (M+H).

Example 64



(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[4-
 20 (dimethylamino)butoxy]benzyl}-2-hydroxypropylcarbamate
 (275)

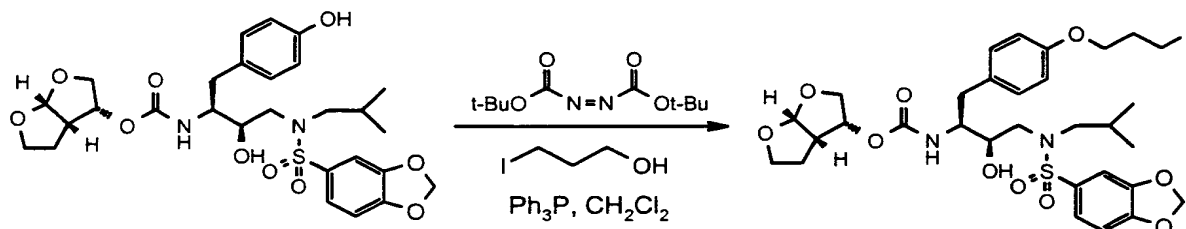
According to example 273, (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-
 b]furan-3-yl (1*S*,2*R*)-1-[4-(4-aminobutoxy)benzyl]-3-[(1,3-
 benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-

25 hydroxypropylcarbamate was subjected to reductive
 alkylation with formaldehyde to give the desired compound
 as a white foam in 74% yield. ¹H NMR (CDCl₃): 7.37 (d,
 1H), 7.22 (s, 1H), 7.15 (d, 2H), 6.93 (d, 1H), 6.86 (d,
 2H), 6.12 (s, 2H), 5.69 (d, 1H), 5.07 (q, 1H), 4.94 (d,
 30 1H), 4.06-3.60 (m, 9H), 3.17 (dd, 1H), 3.00 (m, 4H), 2.81

(m, 2H), 2.40 (t, 2H), 2.32 (s, 6H), 1.95-1.50 (m, 7H), 0.98 (d, 3H), 0.91 (d, 3H). MS(ESI): 692(M+H).

5

Example 65



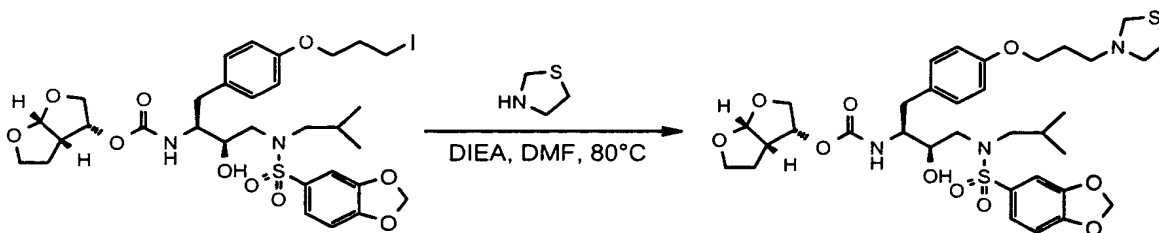
(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-

10 hydroxy-1-[4-(3-iodopropoxy)benzyl]propylcarbamate (276)

The title compound was prepared according to example 202
with the exception that 3-iodo-1-propanol was used
instead of phenethyl alcohol. ¹H NMR (CDCl₃): 7.29 (d,
1H), 7.13 (s, 1H), 7.09 (d, 2H), 6.86 (d, 1H), 6.78 (d,
15 2H), 6.04 (s, 2H), 5.61 (d, 1H), 4.99 (q, 1H), 4.85 (d,
1H), 3.94 (m, 3H), 3.80 (m, 3H), 3.67 (m, 2H), 3.55 (m,
1H), 3.32 (t, 2H), 3.09 (dd, 1H), 3.92 (m, 4H), 2.73 (m,
2H), 2.21 (m, 2H), 1.80 (m, 1H), 1.63 (m, 1H), 1.49 (m,
1H), 0.90 (d, 3H), 0.83 (d, 3H). MS(ESI): 761(M+H).

20

Example 66

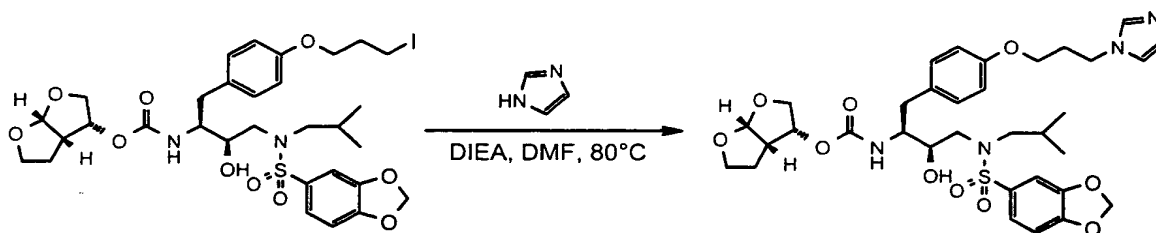


25

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-[3-(1,3-thiazolidin-3-yl)propoxy]
 benzyl]propylcarbamate (277)

- 5 A solution of 35 mg (0.046 mmol) of (3*R*,3*aS*,6*aR*)-
 hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-
 benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-[4-
 (3-iodopropoxy)benzyl]propylcarbamate, 11 μ L (0.14 mmol)
 of thiazolidine, and 32 μ L (0.18 mmol) of *N,N*-
 10 diisopropylethylamine in 0.5 mL of anhydrous DMF was
 heated to 80°C with stirring. After 1.5 hours TLC
 indicated the reaction to be complete. The solution was
 cooled to RT and concentrated *in vacuo*. The residue was
 subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/MeOH)
 15 to afford 22 mg (67%) of the desired compound as a white
 foam. ¹H NMR (CDCl₃): 7.29 (d, 1H), 7.13 (s, 1H), 7.08
 (d, 2H), 7.85 (d, 1H), 6.77 (d, 2H), 6.04 (s, 2H), 5.60
 (d, 1H), 5.00 (q, 1H), 4.88 (d, 1H), 4.01 (s, 2H), 4.00-
 3.88 (m, 3H), 3.80 (m, 3H), 3.67 (m, 2H), 3.55 (br s,
 20 1H), 3.19-3.01 (m, 4H), 3.00-2.65 (m, 7H), 2.52 (t, 2H),
 1.93 (m, 2H), 1.79 (m, 1H), 1.70-1.42 (m, 2H), 0.89 (d,
 3H), 0.83 (d, 3H). MS(ESI): 722 (M+H).

Example 67

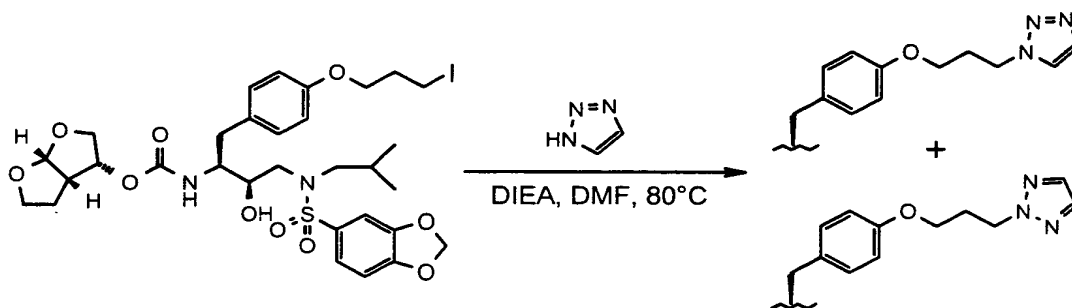


(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-

hydroxy-1-{4-[3-(1*H*-imidazol-1-yl)propoxy]benzyl}
propylcarbamate (278)

The title compound was prepared according to example 66
with the exception that imidazole was substituted for
5 thiazolidine. ¹H NMR (CDCl₃): 7.54 (s, 1H), 7.38 (d, 1H),
7.21 (s, 1H), 7.17 (d, 2H), 7.11 (s, 1H), 6.92 (m, 2H),
6.82 (d, 2H), 6.13 (s, 2H), 5.70 (d, 1H), 5.09 (m, 2H),
4.22 (t, 2H), 4.07-3.68 (m, 8H), 3.22-2.73 (m, 8H), 2.25
(m, 2H), 1.88 (m, 1H), 1.77-1.55 (m, 2H), 0.93 (m, 6H).
10 MS (ESI): 701 (M+H).

Example 68



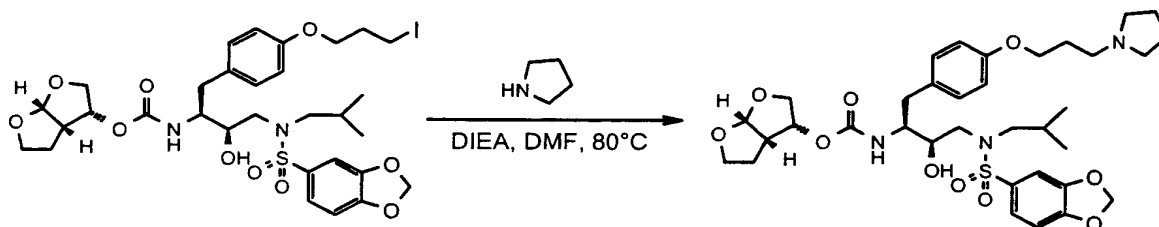
(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-{4-[3-(1*H*-1,2,3-triazol-1-
yl)propoxy]benzyl}propylcarbamate and (3*R*,3*aS*,6*aR*)-
15 hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-
benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-{4-
20 [3-(2*H*-1,2,3-triazol-2-yl)propoxy]benzyl}propylcarbamate
(279)

According to example 66, (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-
25 *b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-
ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-[4-(3-
iodopropoxy)benzyl]propylcarbamate was reacted with

1,2,3-triazole to afford an approximately 1:1 mixture of triazole isomers (as determined by ¹H-NMR and HPLC).
MS (ESI): 702 (M+H).

5

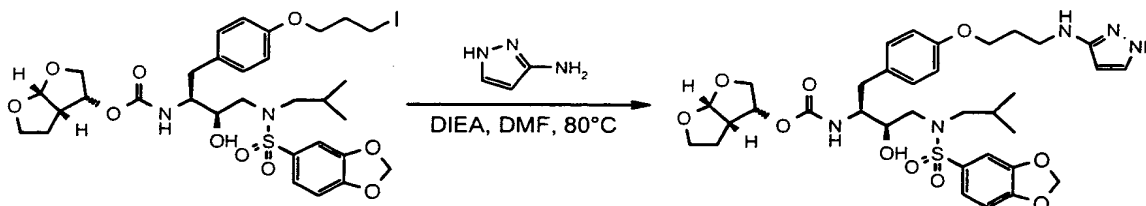
Example 69



(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
10 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-{4-[3-(1-pyrrolidinyl)propoxy]benzyl}
propylcarbamate (280)

The title compound was prepared according to example 66
with the exception that pyrrolidine was substituted for
15 thiazolidine. ¹H NMR (CDCl₃): 7.36 (d, 1H), 7.21 (s, 1H),
7.13 (d, 2H), 6.92 (d, 1H), 6.83 (d, 2H), 6.12 (s, 2H),
5.68 (d, 1H), 5.02 (m, 2H), 4.00 (m, 3H), 3.93-3.62 (m,
6H), 3.15 (dd, 1H), 3.00 (m, 4H), 2.88-2.61 (m, 8H), 2.10
(m, 2H), 1.88 (m, 5H), 1.78-1.50 (m, 2H), 0.96 (d, 3H),
20 0.91 (d, 3H). MS (ESI): 704 (M+H).

Example 70

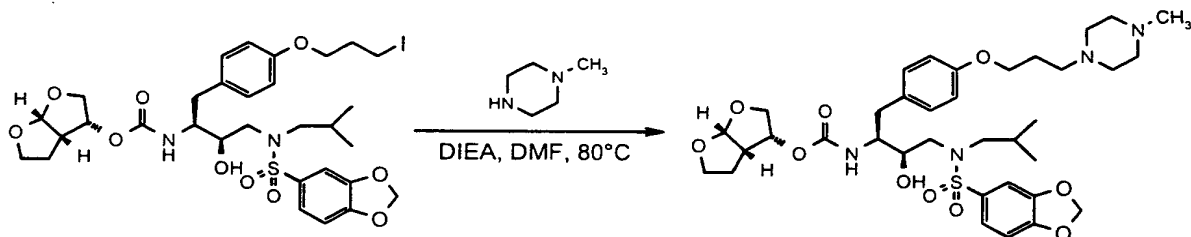


25

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-{4-[3-(1H-pyrazol-3-ylamino)propoxy]benzyl}
 propylcarbamate (281)

5 The title compound was prepared according to example 66
 with the exception that 3-aminopyrazole was substituted
 for thiazolidine. ¹H NMR (CDCl₃): 7.28 (m, 2H), 7.14 (s,
 1H), 7.05 (d, 2H), 6.84 (d, 1H), 6.75 (d, 2H), 6.03 (s,
 2H), 5.58 (m, 2H), 5.15 (d, 1H), 4.99 (q, 1H), 4.80-4.20
 10 (br s, 2H), 4.00 (t, 2H), 3.90 (dd, 1H), 3.78 (m, 3H),
 3.67 (m, 2H), 3.31 (t, 2H), 3.08 (m, 1H), 3.01-2.63 (m,
 7H), 2.01 (m, 2H), 1.80 (m, 1H), 1.61 (m, 1H), 1.47 (m,
 1H), 0.88 (d, 3H), 0.81 (d, 3H). MS(ESI): 716(M+H).

Example 71



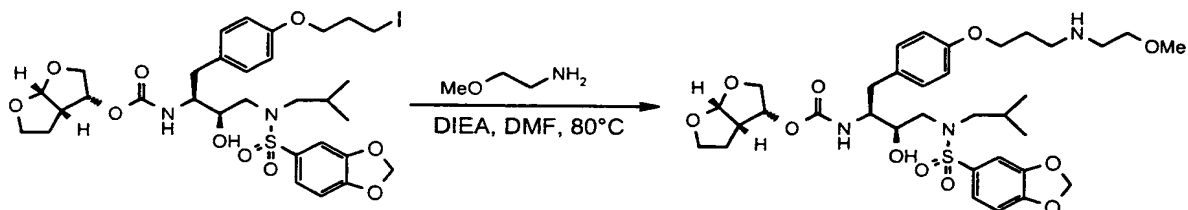
20 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-{4-[3-(4-methyl-1-piperazinyl)propoxy]
 benzyl}propylcarbamate (282)

The title compound was prepared according to example 66
 25 with the exception that 1-methylpiperazine was
 substituted for thiazolidine. ¹H NMR (CDCl₃): 7.37 (d,
 1H), 7.20 (s, 1H), 7.14 (d, 2H), 6.92 (d, 1H), 6.84 (d,
 2H), 6.12 (s, 2H), 5.68 (d, 1H), 5.14-4.92 (m, 2H), 4.00
 (m, 3H), 3.86 (m, 3H), 3.72 (m, 3H), 3.16 (dd, 1H), 3.08-

2.90 (m, 4H), 2.89-2.39 (m, 12H), 2.33 (s, 3H), 2.00 (m, 2H), 1.87 (m, 1H), 1.68 (m, 1H), 1.52 (m, 1H), 0.98 (d, 3H), 0.91 (d, 3H). MS(ESI): 733 (M+H).

5

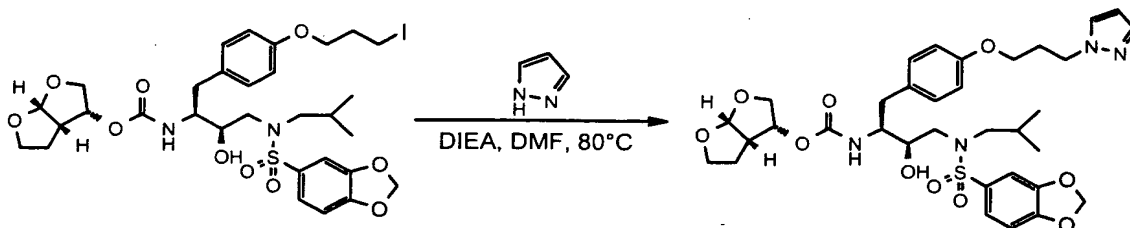
Example 72



10 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-(4-{3-[(2-methoxyethyl)amino]propoxy}benzyl)
propylcarbamate (283)

The title compound was prepared according to example 66
15 with the exception that 2-methoxyethylamine was
substituted for thiazolidine. ¹H NMR (CDCl₃): 7.38 (d,
1H), 7.20 (s, 1H), 7.14 (d, 2H), 6.91 (d, 1H), 6.86 (d,
2H), 6.12 (s, 2H), 5.69 (d, 1H), 5.13-4.90 (m, 2H), 4.01
(m, 3H), 3.87 (m, 3H), 3.74 (m, 3H), 3.52 (t, 2H), 3.40
20 (s, 3H), 3.18 (dd, 1H), 3.00 (m, 4H), 2.81 (m, 7H), 2.08-
1.50 (m, 5H), 0.98 (d, 3H), 0.91 (d, 3H). MS(ESI):
708 (M+H).

Example 73



25

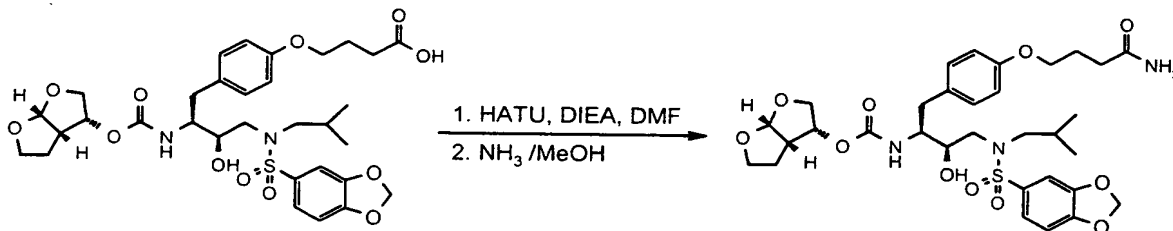
(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-[4-[3-(1*H*-pyrazol-1-yl)propoxy]benzyl]

5 **propylcarbamate (284)**

The title compound was prepared according to example 66 with the exception that pyrazole was substituted for thiazolidine. ¹H NMR (CDCl₃): 7.50 (s, 1H), 7.32 (s, 1H), 7.29 (d, 1H), 7.13 (s, 1H), 7.08 (d, 2H), 6.85 (d, 1H), 6.76 (d, 2H), 6.20 (s, 1H), 6.06 (s, 2H), 5.61 (d, 1H), 5.00 (q, 1H), 4.88 (d, 1H), 4.33 (t, 2H), 3.92 (dd, 1H), 3.80 (m, 5H), 3.68 (m, 2H), 3.58 (br s, 1H), 3.09 (dd, 1H), 2.90 (m, 4H), 2.73 (m, 2H), 2.28 (m, 2H), 1.85-1.43 (m, 3H), 0.89 (d, 3H), 0.82 (d, 3H). MS(ESI): 701(M+H).

15

Example 74



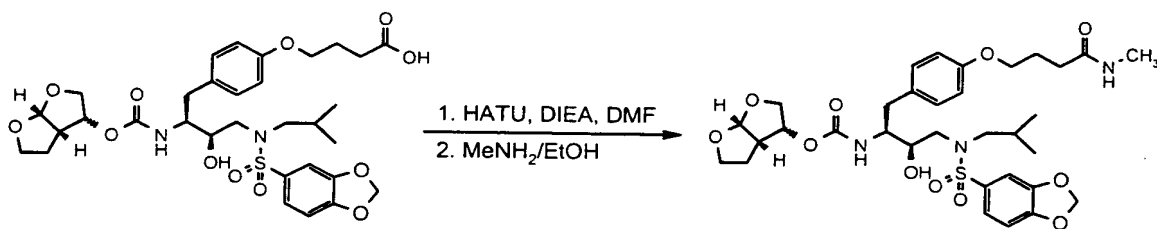
20 **(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-(4-amino-4-oxobutoxy)benzyl]-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate (285)**

A solution of 25 mg (0.037 mmol) of 4-(4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenoxy)butanoic acid and 19 μ L (0.11 mmol) of *N,N*-diisopropylethylamine in 1 mL of anhydrous DMF was

25

treated with 28 mg (0.074 mmol) of O-(7-azabenzotriazol-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate (HATU). The resulting solution was stirred at RT for 15 minutes and was then treated with 0.5 mL of 2M NH₃ in MeOH. After 20 minutes TLC indicated the reaction to be complete. The solution was concentrated to dryness at reduced pressure and the residue subjected to flash chromatography (SiO₂, 95:5 CH₂Cl₂/2M NH₃ in MeOH) to afford 11 mg (44%) of the desired compound as a white powder. ¹H NMR (CDCl₃): 7.30 (d, 1H), 7.14 (s, 1H), 7.07 (d, 2H), 6.85 (d, 2H), 6.75 (d, 2H), 6.05 (s, 2H), 5.60 (d, 1H), 5.56-5.34 (m, 2H), 4.97 (m, 2H), 3.94 (m, 3H), 3.78 (m, 3H), 3.66 (m, 2H), 3.09 (m, 1H), 3.00-2.64 (m, 7H), 2.40 (t, 2H), 2.08 (m, 2H), 1.80 (m, 1H), 1.61 (m, 1H), 1.48 (m, 1H), 0.90 (d, 3H), 0.82 (d, 3H). MS(ESI): 678 (M+H).

Example 75



20

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-{4-[4-(methylamino)-4-oxobutoxy]benzyl}propylcarbamate (286)

25

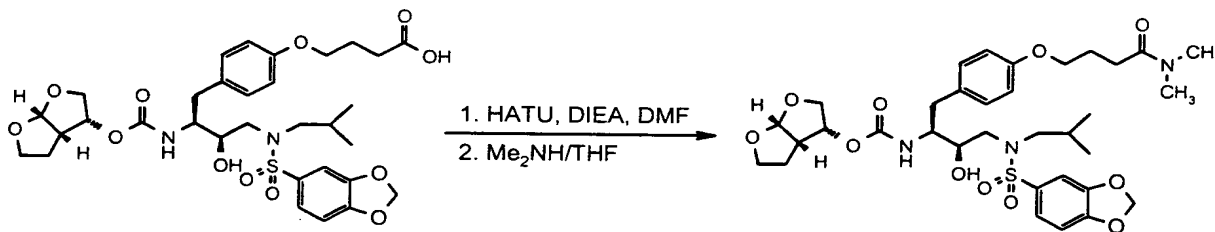
A solution of 35 mg (0.052 mmol) of 4-(4-{(2S,3R)-2-([[(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-3-

hydroxybutyl}phenoxy)butanoic acid and 27 μ L (0.16 mmol) of *N,N*-diisopropylethylamine in 1 mL of anhydrous DMF was treated with 49 mg (0.13 mmol) of *O*-(7-azabenzotriazol-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate

5 (HATU). The resulting solution was stirred at RT for 15 minutes and was then treated with 0.5 mL of 8M MeNH₂ in EtOH. After 18 hours TLC indicated the reaction to be complete. The solution was concentrated to dryness at

10 chromatography (SiO₂, 95:5 CH₂Cl₂/MeOH) to afford 22 mg (61%) of the desired compound as a white powder. ¹H NMR (CDCl₃): 7.37 (d, 1H), 7.22 (s, 1H), 7.13 (d, 2H), 6.92 (d, 1H), 6.82 (d, 2H), 6.13 (s, 2H), 5.67 (m, 2H), 5.05 (m, 2H), 4.00 (m, 3H), 3.88 (m, 3H), 3.72 (m, 2H), 3.24-
15 2.71 (m, 11H), 2.40 (t, 2H), 2.16 (m, 2H), 1.88 (m, 1H), 1.67 (m, 1H), 1.53 (m, 1H), 0.94 (m, 6H). MS(ESI): 692 (M+H).

Example 76



(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[4-
(dimethylamino)-4-oxobutoxy]benzyl}-2-

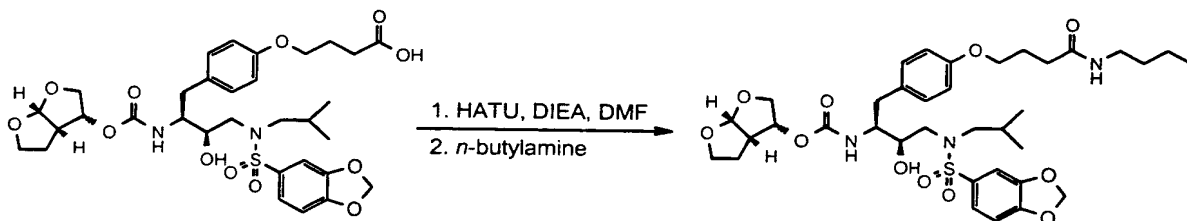
25 hydroxypropylcarbamate (287)

The title compound was prepared according to example 286 with the exception that 1 mL of 2M Me₂NH/THF was substituted for MeNH₂/EtOH. ¹H NMR (CDCl₃): 7.36 (d, 1H), 7.21 (s, 1H), 7.13 (d, 2H), 0.92 (d, 1H), 6.85 (d, 2H),

6.12 (s, 2H), 5.69 (d, 1H), 5.02 (m, 2H), 4.01 (m, 3H), 3.87 (m, 3H), 3.73 (m, 2H), 3.16 (dd, 1H), 3.10-2.88 (m, 11H), 2.81 (m, 2H), 2.54 (t, 2H), 2.16 (m, 2H), 1.86 (m, 1H), 1.79-1.50 (m, 2H), 0.98 (d, 3H), 0.90 (d, 3H).

5 MS(ESI): 706(M+H).

Example 77

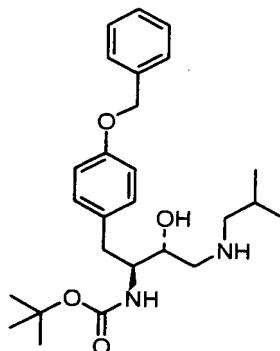


10 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[4-
(butylamino)-4-oxobutoxy]benzyl}-2-hydroxypropylcarbamate
(288)

The title compound was prepared according to example 286
15 with the exception that 60 μ L of *n*-butylamine was
substituted for $\text{MeNH}_2/\text{EtOH}$. ^1H NMR (CDCl_3): 7.38 (d, 1H),
7.21 (s, 1H), 7.14 (d, 2H), 6.93 (d, 1H), 6.84 (d, 2H),
6.12 (s, 2H), 5.69 (d, 1H), 5.60 (br s, 1H), 5.03 (m,
2H), 4.00 (m, 3H), 3.87 (m, 3H), 3.73 (m, 2H), 3.39-2.72
20 (m, 10H), 2.39 (t, 2H), 2.13 (m, 2H), 1.88 (m, 1H), 1.69
(m, 1H), 1.60-1.27 (m, 5H), 0.93 (m, 9H). MS(ESI):
734 (M+H).

Example 78

25 Step 1:
t-Butyl-(1S,2R)-1-(4-benzyloxy-benzyl)-3-*i*-butylamino-2-
hydroxypropyl-carbamate

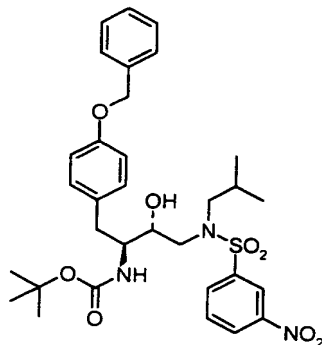


5 *i*-Butylamine (7.0 g, 94.7 mmol) was added to a solution
of *t*-butyl-(1*S*,2*R*)-1-(4-benzyloxy-benzyl)-2,3-epoxydo-
propyl-carbamate (5.0 g, 13.5 mmol) (prepared according
to the reference by Chen, P. *at all.*, *Tetrahedron Letters*
1997, 38, 3175-3178), in *i*-propanol (70 mL). The mixture
10 was stirred at 85°C for 2 hours, then cooled to 5°C. Water
(400 mL) was added dropwise. The resulting suspension was
stirred for 30 minutes at 5°C, then filtered. The solid
was washed with water (3x100 mL), and dried *in vacuo* to
yield title product (5.5 g, 92%).

15 ¹H NMR (CDCl₃): δ 0.88 (6H, d), 1.34 (9H, s), 1.68 (1H,
m), 2.38 (2H, d), 2.64 (2H, d), 2.80 (1H, m), 2.86 (1H,
dd), 3.40 (1H, m), 3.70 (2H, broad s), 4.63 (1H, d), 5.01
(2H, s), 6.88 (2H, d), 7.12 (2H, d), 7.40 (5H, m).

20 Step 2:

t-Butyl-N((1*S*,2*R*)-1-(4-benzyloxy-benzyl)-3-*i*-butyl-[(3-nitrobenzene)sulfonyl]-amino-2-hydroxypropyl-carbamate

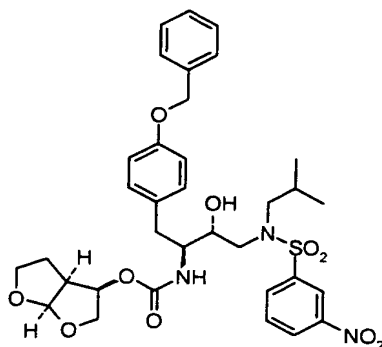


5 A CH₂Cl₂ (50 mL) solution of the product from Step 1 (1.7 g, 3.8 mmol), 3-nitrobenzenesulfonylchloride (1.1 g, 4.8 mmol), and diisopropylethylamine (1.0 g, 7.8 mmol) was stirred at 20°C for 4 hours. Water (70 mL) was then added to the reaction mixture and the phases were separated. The aqueous phase was extracted with CH₂Cl₂ (3x100 mL).
10 The combined organic phases were dried over MgSO₄ and concentrated. The residue was dissolved in ether (300 mL), and silicagel (50 g) was added to the solution. The mixture was then filtered. The filtrate was concentrated under reduced pressure. The residue was added hexane (300 mL), stirred for three hours at 20°C, and then the white solid was filtered and dried to afford title compound (1.9 g, 80%).

¹H NMR (CDCl₃): δ 0.88 (6H, d), 1.35 (9H, s), 1.86 (1H, m), 2.85 (2H, m), 2.99 (2H, d), 3.19 (2H, d), 3.77 (2H, m), 4.57 (1H, d), 5.03 (2H, s), 6.90 (1H, d), 7.12 (1H, d), 7.40 (5H, m), 7.69 (1H, t), 8.08 (1H, d), 8.39 (1H, d), 8.62 (1H, s).

25 Step 3:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-benzyloxy-benzyl)-3-*i*-butyl-[(3-nitrobenzene)sulfonyl]-amino-2-hydroxypropyl)-carbamate



5

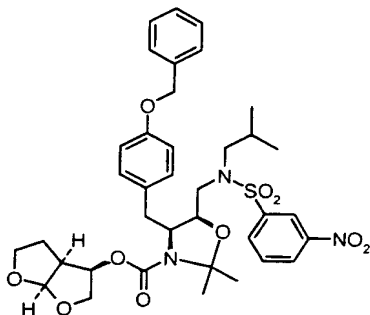
Trifluoroacetic acid (15 mL) was added dropwise to a solution of the product from Step 2 (1.5 g, 5 mmol) in CH₂Cl₂ (40 mL). The mixture was stirred for 1 hour at 20°C, then concentrated under reduced pressure. The residue was dissolved in CH₂Cl₂ (200 mL) and 10% aqueous Na₂CO₃ solution (100 mL) was added. The mixture was stirred at 20°C for 15 minutes. Phases were separated, and the aqueous phase was extracted with additional CH₂Cl₂ (2x50 mL). The combined organic phases were then dried over Na₂CO₃ and concentrated under reduced pressure. The residue was dissolved in acetonitrile (40 mL). Diisopropylethylamine (2.0 g, 15.3 mmol) and (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl-4-nitrophenyl carbonate (2.0 g, 6.7 mmol) were added, and the solution was stirred for 12 hours at 20°C. Then 25% aqueous ammonium hydroxide solution (6 mL) was added, and the mixture was stirred for an additional 0.5 hour. The solvents were removed under reduced pressure. The residue was dissolved in EtOAc (100 mL), and the solution was

extracted with 5% aqueous Na_2CO_3 solution (10x75 ml) and washed with brine (1x75 mL). The organic phase was dried over Na_2CO_3 , and concentrated in vacuo. Hexane (30 mL) was added to the residue and the mixture was stirred at 20°C for 1 hour. Then the white solid was filtered and air dried to yield title compound (1.5 g, 60%).

^1H NMR (CDCl_3): δ 0.86 (3H, d), 0.88 (3H, d), 1.54 (1H, m), 1.65 (1H, m), 1.85 (1H, m), 2.75 (1H, m), 2.97 (4H, m), 3.14 (2H, m), 3.37 (1H, m), 3.68 (2H, m), 3.82 (2H, m), 3.94 (2H, m), 4.85 (1H, d), 5.00 (3H, s), 5.60 (1H, d), 6.88 (2H, d), 7.09 (2H, d), 7.35 (5H, m), 7.72 (1H, t), 8.07 (1H, d), 8.38 (1H, d), 8.61 (1H, s); MS: 684 (M^+).

Step 4:

N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-(4S,5R)-4-(4-benzyloxy-benzyl)-5-*i*-butyl-[(3-nitrobenzene)sulfonyl]-aminomethyl-2,2-dimethyl-oxazolidine



A CH_2Cl_2 (8 mL) solution of the product of Step 3 (0.5 g, 0.7 mmol) was added 2,2-dimethoxy-propane (0.8 mL, 6.5 mmol) and *p*-toluenesulfonic acid (40 mg, 0.2 mmol). The mixture was refluxed for 4 hours, cooled to 20°C, and washed with 5% aqueous Na_2CO_3 solution (10 mL). The

organic phase was dried over MgSO_4 , filtered, and concentrated under reduced pressure. The residue was chromatographed on SiO_2 using hexane-EtOAc (1:1) as eluent, to provide title compound (0.5 g, 95%).

5

^1H NMR (CDCl_3): δ 0.83 (3H, d), 0.91 (3H, d), 1.33, 1.40 (3H, s)*, 1.50, 1.57 (3H, s)*, 1.85 (2H, m), 1.97 (1H, m), 2.70 (3H, m), 3.10 (3H, m), 3.22 (1H, d), 3.36 (1H, d), 3.43 (1H, m), 3.80 (2H, m), 3.94 (1H, m), 4.21 (2H, m), 5.03 (2H, s), 5.65, 5.68 (1H, d)*, 6.87 (2H, d), 7.02 (1H, d), 7.38 (5H, m), 7.62 (1H, t), 7.85 (1H, d), 8.35 (1H, t), 8.54 (1H, s); MS: 724 (M^+).

10

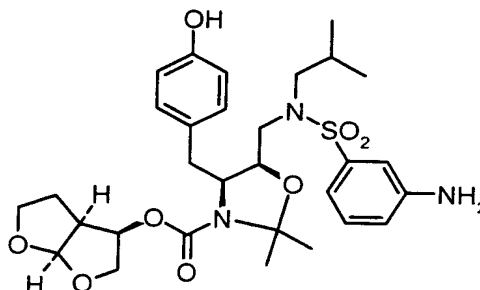
*possible indication for rotamers.

15

Step 5:

N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-(4S,5R)-4-(4-hydroxybenzyl)-5-*i*-butyl-[(3-aminobenzene)sulfonyl]-aminomethyl-2,2-dimethyl-oxazolidine

20



Pd/C (10% Pd, Degussa type) (0.5 g) was added to a THF (10 mL) solution of the product of Step 4 (0.5 g; 0.7 mmol). The mixture was hydrogenated under atmospheric pressure H_2 for 12 hours. The catalyst was removed by

25

filtration through Celite, and the solvent was evaporated in vacuo. The residue was triturated in hexane. The white solid was then filtered and washed with hexane (2x20 ml) to afford title compound (210 mg, 50%).

5

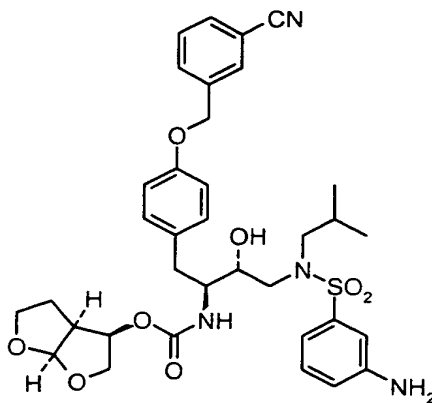
^1H NMR ($\text{DMSO}-d_6$): δ 0.80 (6H, m), 1.22, 1.35 (3H, s)*, 1.45, 1.53 (3H, s)*, 1.75 (2H, m), 1.95 (1H, m), 2.50-3.30 (10H, m), 3.60 (2H, d), 3.80 (1H, m), 4.06 (1H, m), 4.74 (1H, dd), 5.52 (2H, d), 6.65 (3H, m), 6.90 (3H, m), 7.12 (1H, t), 9.22 (1H, s); MS: 604 (M^+).

10

*possible indication for rotamers.

Step 6:

15 (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-[4-(3-cyanobenzyloxy)-benzyl]-3-*i*-butyl-[(3-aminobenzene)sulfonyl]-amino-2-hydroxypropyl)-carbamate (289)



20

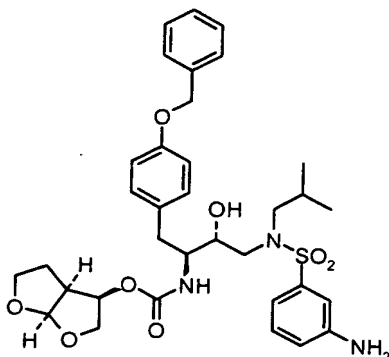
Cs_2CO_3 (134.4 mg, 0.4 mmol) and 3-(bromomethyl)benzonitrile (65 mg, 0.3 mmol) were added to the product of Step 5 (200 mg, 0.3 mmol) in DMF (2 mL). The mixture was stirred at 20°C for 4 hours, diluted with

Et₂O (50 mL), and filtered. The filtrate was washed with water (1x35 mL), dried over MgSO₄, and concentrated under reduced pressure. The residue, dissolved in 1,4-dioxane (6 mL), was added 4 M HCl in dioxane (7.5 ml) and water (0.2 g). The solution was stirred at 20°C for 2 hours and then diluted with water (50 mL). A solution of 5% aqueous Na₂CO₃ was added until pH 12-14, and the mixture was extracted with CH₂Cl₂ (3x20 mL). The combined organic phases were dried over MgSO₄ and concentrated under reduced pressure. The residue was chromatographed on SiO₂ using EtOAc-hexane (4:1) as eluent to afford title compound (85 mg, 45%).

¹H NMR (DMSO-d₆): δ 0.78 (3H, d), 0.83 (3H, d), 1.25 (2H, m), 1.34 (1H, m), 1.90 (1H, m), 2.36 (1H, t), 2.71 (3H, m), 2.93 (2H, m), 3.54 (4H, m), 3.66 (1H, t), 3.80 (1H, dd), 4.80 (1H, dd), 4.95 (1H, d), 5.08 (2H, s), 5.47 (1H, d), 5.53 (2H, s), 6.73 (1H, d), 6.80 (1H, d), 6.84 (2H, d), 6.92 (1H, s), 7.10 (2H, d), 7.15 (2H, t), 7.57 (1H, t), 7.75 (2H, t), 7.85 (1H, s); MS: 679 (M⁺).

Example 79

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-benzyloxy-benzyl)-3-*i*-butyl-[(3-aminobenzene)sulfonyl]-amino-2-hydroxypropyl)-carbamate (290)

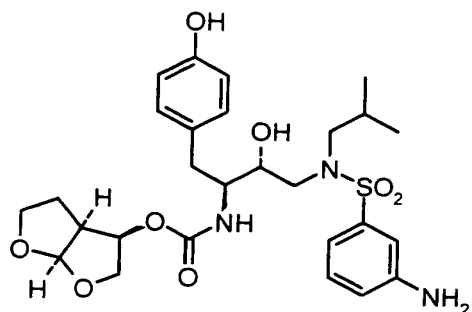


(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-benzyloxy-benzyl)-3-i-butyl-[(3-nitrobenzene)sulfonyl]-amino-2-hydroxypropyl)-carbamate (100 mg, 0.15 mmol) in THF (3 mL) was added Pt/C (3% Pt) (30 mg). The mixture was hydrogenated under atmospheric pressure H₂ for 12 hours. The catalyst was removed by filtration through Celite, and the solvent was evaporated in vacuo to yield title compound (60 mg, 65%).

¹H NMR (CDCl₃): δ 0.82 (3H, d), 0.89 (3H, d), 1.22 (1H, m), 1.60 (1H, m), 1.81 (2H, m), 2.77 (2H, m), 2.97 (4H, m), 3.15 (1H, m), 3.64 (3H, m), 3.82 (3H, m), 3.95 (1H, m), 4.96 (1H, d), 5.02 (3H, d), 5.63 (1H, d), 6.80 (1H, d), 6.88 (2H, d), 6.96 (3H, m), 7.08 (3H, m), 7.12 (1H, m), 7.35 (4H, m); MS: 654 (M⁺).

Example 80

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-hydroxybenzyl)-3-i-butyl-[(3-aminobenzene)sulfonyl]-amino-2-hydroxypropyl)-carbamate (291)

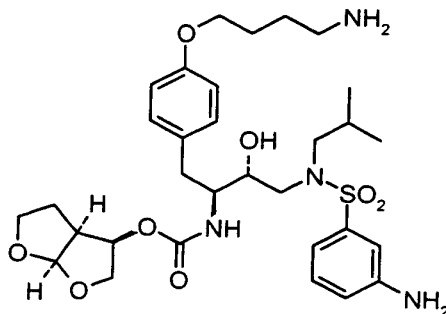


(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-
 5 (4-benzyloxy-benzyl)-3-*i*-butyl-[(3-nitrobenzene)sulfonyl]-amino-2-hydroxypropyl)-carbamate
 (300 mg, 0.44 mmol) in THF (8 mL) was added Pd/C (10% Pd,
 Degussa Type) (300 mg). The mixture was hydrogenated
 under atmospheric pressure H₂ for 12 hours. The catalyst
 10 was removed by filtration through Celite, and the solvent
 was evaporated in vacuo to yield title compound (95 mg,
 50%).

¹H NMR (DMSO-*d*₆): δ 0.75 (3H, d), 0.82 (3H, d), 1.25 (2H,
 15 m), 1.40 (1H, m), 1.93 (1H, m), 2.32 (1H, t), 2.72 (3H,
 m), 2.85 (1H, d), 2.95 (1H, m), 3.55 (4H, m), 3.71 (1H,
 t), 3.81 (1H, dd), 4.82 (1H, dd), 4.92 (1H, m), 5.48 (1H,
 d), 5.53 (2H, s), 6.56 (2H, d), 6.73 (1H, d), 6.79 (1H,
 d), 6.94 (3H, m), 7.14 (2H, m), 9.00 (1H, s).

Example 81

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-(butylamine)benzyl)-3-*i*-butyl-[(3-aminobenzene)sulfonyl]-amino-2-hydroxypropyl-carbamate (292)



5

t-Butyl-N-(4-hydroxybutyl)carbamate (75.7 mg, 0.40 mmol) and triphenylphosphine (105 mg, 0.40 mmol) were mixed in CH₂Cl₂ (5 mL) and stirred at 0°C for 30 minutes. Di-t-butyl azodicarboxylate (92.1 mg, 0.40 mmol) was then added, followed by (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-hydroxybenzyl)-3-*i*-butyl-[(3-aminobenzene)sulfonyl]-amino-2-hydroxypropyl-carbamate (90 mg, 0.16 mmol). The mixture was stirred at 20°C for 12 hours. The solvent was then evaporated, and the residue was chromatographed on SiO₂ using EtOAc-hexane (4:1) as eluent. The isolated intermediate was dissolved in CH₂Cl₂ (16 mL). TFA (4 mL) was added. The mixture was stirred for 2 hours at room temperature, and concentrated to afford title compound (65 mg, 50%) as a TFA salt.

20

¹H NMR (DMSO-*d*₆): δ 0.75 (3H, d), 0.82 (3H, d), 1.20 (2H, m), 1.38 (1H, m), 1.60 (6H, m), 1.90 (1H, m), 2.38 (1H, t), 2.75 (3H, m), 2.92 (1H, m), 3.37 (2H, m), 3.72 (1H, t), 3.82 (1H, dd), 3.90 (1H, m), 4.37 (1H, t), 4.80 (1H, dd), 5.48 (1H, d), 6.77 (3H, m), 6.85 (1H, d), 6.97 (1H,

25

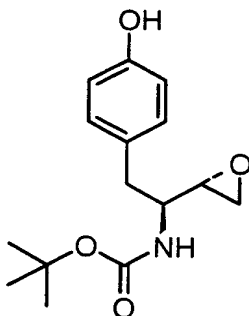
s), 7.09 (2H, d), 7.16 (2H, dd), 7.65 (2H, broad s); MS: 635 (M⁺).

5

Example 82

Step 1:.

t-Butyl-(1S,2R)-1-(4-hydroxybenzyl)-2,3-epoxydo-propyl-carbamate



10

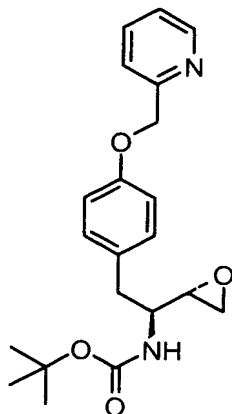
Pd(OH)₂/C (20% Pd, Degussa Type) (70 mg) was added to a solution of t-butyl-(1S,2R)-1-(4-benzyloxy-benzyl)-2,3-epoxydo-propyl-carbamate (700 mg, 1.9 mmol) (prepared according to the reference by Chen, P. at all., *Tetrahedron Letters* 1997, 38, 3175-3178) in EtOH (12 mL) and EtOAc (3 mL). The mixture was hydrogenated under atmospheric pressure H₂ for 2 hours, filtrated through Celite, and concentrated to yield title compound (530 mg, 100%).

20

¹H NMR (DMSO-d₆): δ 1.25 (9H, s), 2.55 (4H, m), 2.82 (1H, m), 3.36 (1H, m), 6.60 (2H, d), 6.75 (1H, d), 6.95 (2H, d), 9.09 (1H, s).

25 **Step 2:**

t-Butyl-(1*S*,2*R*)-1-(4-(2-pyridylmethoxy)benzyl)-2,3-epoxydo-propyl-carbamate



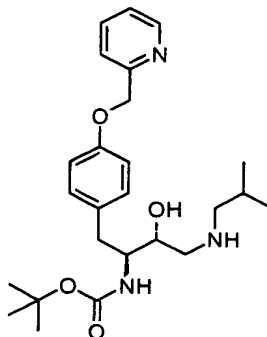
5 Cs₂CO₃ (1.60 g, 5.05 mmol) in DMF (10 mL) was added 2-picolylchloride hydrochloride (400 mg, 2.44 mmol). After stirring for 5 minutes at room temperature, *t*-Butyl-(1*S*,2*R*)-1-(4-hydroxybenzyl)-2,3-epoxydo-propyl-carbamate (450 mg, 1.61 mmol) was added. The mixture was stirred
10 for an additional 2 hours at 20°C, diluted with Et₂O (50 mL), and filtered. The filtrate was washed with water, dried over MgSO₄, and concentrated. The residue was chromatographed on silicagel using EtOAc-hexane (3:2) as eluent to afford title compound (505 mg, 85%).

15

¹H NMR (DMSO-*d*₆): δ 1.22 (9H, s), 2.45 (1H, s), 2.62 (1H, m), 2.74 (1H, dd), 2.84 (1H, m), 3.42 (1H, m), 5.08 (2H, s), 6.79 (1H, d), 6.87 (2H, d), 7.07 (2H, d), 7.28 (1H, dd), 7.44 (1H, d), 7.78 (1H, t), 8.52 (1H, d); MS: 371
20 (M⁺).

Step 3:

***t*-Butyl-(1*S*,2*R*)-1-(4-(2-pyridylmethyloxy)benzyl)-3-*i*-butylamino-2-hydroxypropyl-carbamate**

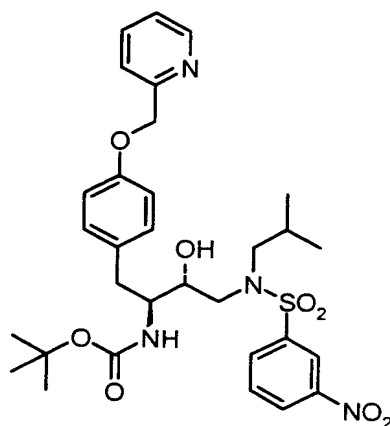


i-Butylamine (610 mg, 8.33 mmol) was added to a solution of *t*-Butyl-(1*S*,2*R*)-1-(4-(2-pyridylmethyloxy)benzyl)-2,3-epoxydo-propyl-carbamate (440 mg, 1.20 mmol) in *i*-propanol (10 mL). The mixture was then stirred at 85°C for 2 hours, cooled to 5°C, diluted with water (75 mL), and extracted with CHCl₃ (4x70 mL). The combined organic phases were dried over MgSO₄, filtered, and concentrated to afford title product (320 mg, 60%).

¹H NMR (CDCl₃): δ 0.87 (6H, d), 1.33 (9H, s), 1.67 (1H, m), 2.37 (2H, d), 2.65 (2H, d), 2.77 (1H, m), 2.85 (1H, dd), 3.41 (1H, m), 3.72 (2H, m), 4.63 (1H, d), 5.15 (2H, s), 6.89 (2H, d), 7.12 (2H, d), 7.19 (1H, dd), 7.49 (1H, d), 7.67 (1H, t), 8.57 (1H, d).

Step 4:

***t*-Butyl-N((1*S*,2*R*)-1-(4-(2-pyridylmethyloxy)benzyl)-3-*i*-butyl-[(3-nitrobenzene)sulfonyl]-amino-2-hydroxypropyl-carbamate**



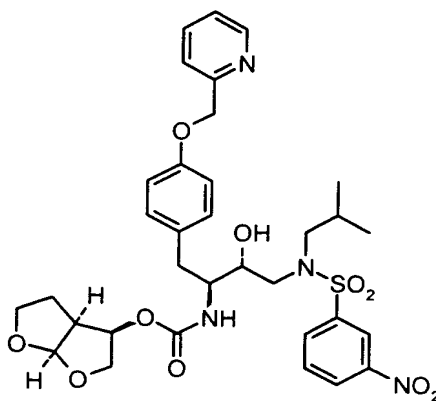
5

A CH₂Cl₂ (10 mL) solution of t-Butyl-(1S,2R)-1-(4-(2-pyridylmethoxy)benzyl)-3-*i*-butylamino-2-hydroxypropyl-carbamate (320 mg, 0.72 mmol), 3-nitrobenzenesulfonyl chloride (200 mg, 0.90 mmol), and diisopropylethylamine (186 mg, 1.44 mmol) was stirred at 20°C for 12 hours. Water (20 mL) was then added to the reaction mixture and the phases were separated. The aqueous phase was extracted with CH₂Cl₂ (3x25 mL). The combined organic phases were dried over MgSO₄ and concentrated. The residue was then filtered through SiO₂ using CH₂Cl₂-acetone (3:2) as eluent. The filtrate was concentrated under reduced pressure to yield title product (330 mg, 75%).

¹H NMR (DMSO-*d*₆): δ 0.78 (6H, d), 1.20 (9H, s), 1.93 (1H, m), 2.37 (1H, dd), 2.85 (1H, dd), 3.10 (2H, m), 3.42 (1H, d), 4.88 (1H, d), 5.08 (2H, s), 6.61 (1H, d), 6.85 (2H, d), 7.04 (2H, d), 7.29 (1H, m), 7.44 (1H, d), 7.82 (2H, m), 8.19 (1H, d), 8.43 (2H, s), 8.53 (1H, d).

5 Step 5:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-(2-pyridylmethoxy)benzyl)-3-*i*-butyl-[(3-nitrobenzene)sulfonyl]-amino-2-hydroxypropyl-carbamate



10

Trifluoroacetic acid (4 mL) was added dropwise to a solution of *t*-Butyl-N((1S,2R)-1-(4-(2-pyridylmethoxy)benzyl)-3-*i*-butyl-[(3-nitrobenzene)sulfonyl]-amino-2-hydroxypropyl-carbamate (330 mg, 0.52 mmol) in CH₂Cl₂ (8 mL). The mixture was stirred at 20°C for 1.5 hour, then concentrated under reduced pressure. The residue was dissolved in CH₂Cl₂ (50 mL) and 10% aqueous Na₂CO₃ solution (50 mL) was added. The mixture was stirred at 20°C for 15 minutes. Phases were separated, and the aqueous phase was extracted with additional CH₂Cl₂ (2x50 ml). The combined organic phases were then dried over Na₂CO₃ and concentrated under reduced

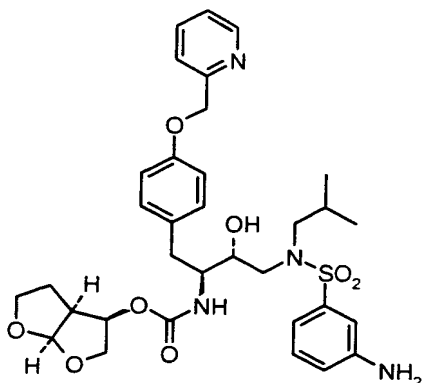
pressure. The residue was dissolved in acetonitrile (7 mL). Diisopropylethylamine (269 mg, 2.08 mmol) and (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl-4-nitrophenyl carbonate (272 mg, 0.92 mmol) were added, and the solution was stirred at 20°C for 12 hours. Then, 25% aqueous ammonium hydroxide solution (2 mL) was added, and the mixture was stirred for an additional 0.5 hour. The solvents were removed under reduced pressure. The residue was dissolved in EtOAc (50 mL), and the solution was extracted with 5% aqueous Na₂CO₃ solution (10x25 mL) and washed with brine (1x75 mL). The organic phase was dried over Na₂CO₃, and concentrated *in vacuo*. The residue was chromatographed on SiO₂ using EtOAc as eluent to afford title compound (175 mg, 50%).

¹H NMR (DMSO-d₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.21 (1H, m), 1.33 (1H, m), 1.92 (1H, m), 2.35 (1H, dd), 2.73 (1H, m), 2.85 (2H, m), 3.05 (1H, m), 3.12 (1H, m), 3.45 (1H, m), 3.52 (1H, m), 3.64 (1H, t), 3.78 (1H, dd), 4.81 (1H, q), 4.97 (1H, d), 5.06 (2H, s), 5.46 (1H, d), 6.85 (2H, d), 7.05 (2H, d), 7.17 (1H, d), 7.29 (1H, dd), 7.44 (1H, d), 7.78 (1H, t), 7.84 (1H, t), 8.19 (1H, d), 8.42 (2H, m), 8.53 (1H, d); MS: 685 (M⁺).

Step 6:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-(2-pyridylmethoxy)benzyl)-3-*i*-butyl-[(3-aminobenzene)sulfonyl]-amino-2-hydroxypropyl-carbamate (293)

30



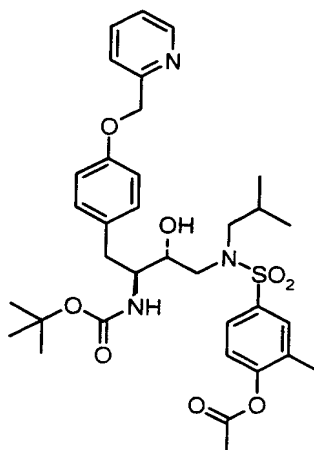
(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-(2-pyridylmethoxy)benzyl)-3-i-butyl-[(3-nitrobenzene)sulfonyl]-amino-2-hydroxypropyl)-carbamate (170 mg, 0.25 mmol) in THF (7 mL) was added Pt/C (3% Pt) (100 mg). The mixture was hydrogenated under atmospheric pressure H₂ for 12 hours. The catalyst was removed by filtration through Celite, and the solvent was evaporated in vacuo. The residue was chromatographed on SiO₂ using EtAOc as eluent to yield title compound (75 mg, 46%).

¹H NMR (DMSO-d₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.21 (1H, m), 1.30 (1H, m), 1.93 (1H, m), 2.36 (1H, t), 2.72 (3H, m), 2.97 (2H, m), 3.55 (4H, m), 3.65 (1H, t), 3.79 (1H, dd), 4.80 (1H, m), 4.95 (1H, d), 5.06 (2H, s), 5.45 (1H, d), 5.53 (2H, broad s), 6.72 (1H, d), 6.79 (1H, d), 6.83 (2H, d), 6.91 (1H, s), 7.08 (2H, d), 7.13 (1H, t), 7.17 (1H, d), 7.28 (1H, d), 7.45 (1H, d), 7.77 (1H, t), 8.53 (1H, d); MS: 655 (M⁺).

Example 83

Step 1:

t-Butyl-N((1*S*,2*R*)-1-(4-(2-pyridylmethyloxy)benzyl)-3-*i*-butyl-[(3-methyl-4-acetoxy)benzenesulfonyl]-amino-2-hydroxypropyl-carbamate

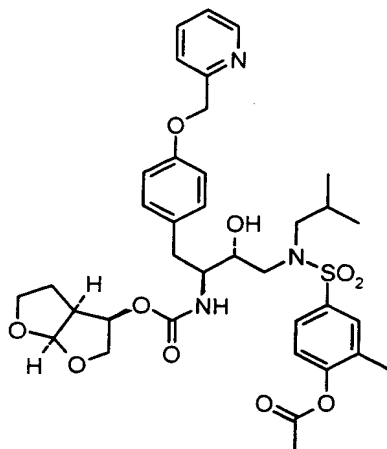


5 A CH₂Cl₂ (15 mL) solution of *t*-Butyl-(1*S*,2*R*)-1-(4-(2-pyridylmethyloxy)benzyl)-3-*i*-butylamino-2-hydroxypropyl-carbamate (630 mg, 1.42 mmol) was added (3-methyl-4-acetoxy)benzenesulfonyl chloride (442 mg, 1.78 mmol), and diisopropylethylamine (367 mg, 2.84 mmol). The mixture
10 was stirred at room temperature for 1 hour, and concentrated. The residue was added water (50 mL), and stirred at 20°C for 30 minutes to afford title compound (675 mg, 75%).

15 MS: 656 (M⁺).

Step 2:

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl-N((1*S*,2*R*)-1-(4-(2-pyridylmethyloxy)benzyl)-3-*i*-butyl-[(3-methyl-4-acetoxy)benzenesulfonyl]-amino-2-hydroxypropyl-carbamate
20 (294)



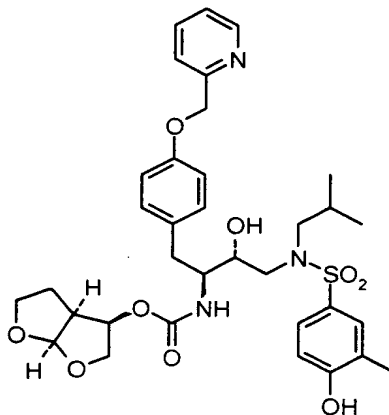
A CH_2Cl_2 (15 mL) and TFA (7.5 mL) solution of *t*-Butyl-
 N((1*S*,2*R*)-1-(4-(2-pyridylmethoxy)benzyl)-3-*i*-butyl-[(3-
 methyl-4-acetoxy)benzenesulfonyl]-amino-2-hydroxypropyl-
 5 carbamate (650 mg, 1 mmol) was stirred at 20°C for 1 hour,
 and concentrated. The residue was dissolved in EtOAc (50
 mL), and washed with 2% aqueous NaHCO_3 solution (3x25 mL).
 The organic phase was dried over MgSO_4 , and filtered.
 Diisopropylethylamine (517 mg, 4 mmol) and (3*R*,3*aS*,6*aR*)-
 10 hexahydrofuro[2,3-*b*]furan-3-yl-4-nitrophenyl carbonate
 (517 mg, 1.75 mmol) were then added, and the solvent was
 evaporated. The residue was dissolved in CH_3CN (20 mL), and
 more diisopropylethylamine (517 mg, 4 mmol) was added.
 The resulting solution was stirred at room temperature
 15 for 3 hours, and concentrated. The residue was dissolved
 in EtOAc (150 mL), then washed with water (1x100 mL), 5%
 aqueous Na_2CO_3 (10x100 mL), and brine (1x100 mL). The
 organic phase was dried over MgSO_4 , filtered, and
 concentrated. The residue was then chromatographed on SiO_2
 20 using EtOAc as eluent to yield title compound (385 mg,
 55%).

¹H NMR (DMSO-*d*₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.20 (1H, m), 1.35 (1H, m), 2.17 (3H, s), 2.30 (3H, s), 2.37 (1H, t), 2.76 (3H, m), 2.91 (1H, d), 3.04 (1H, dd), 3.56 (4H, m), 3.66 (1H, t), 3.79 (1H, dd), 4.80 (1H, dd), 5.01 (1H, d), 5.07 (2H, s), 5.46 (1H, d), 6.85 (2H, d), 7.09 (2H, d), 7.19 (1H, d), 7.26 (1H, d), 7.30 (2H, t), 7.46 (1H, d), 7.61 (1H, d), 7.70 (1H, s), 7.78 (1H, t), 8.53 (1H, d); MS: 712 (M⁺).

10

Example 84

(3R,3aS,6aR)-Hexahydrofuro[2,3-*b*]furan-3-yl-N((1S,2R)-1-(4-(2-pyridylmethyloxy)benzyl)-3-*i*-butyl-[(3-methyl-4-hydroxy)benzenesulfonyl]-amino-2-hydroxypropyl-carbamate (295)



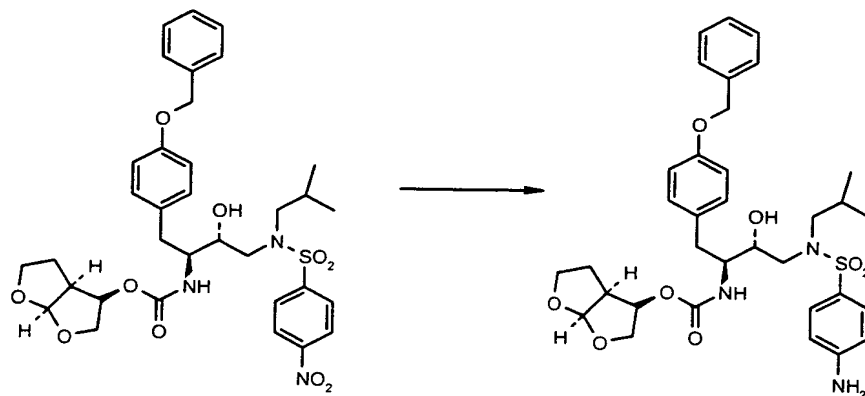
(3R,3aS,6aR)-Hexahydrofuro[2,3-*b*]furan-3-yl-N((1S,2R)-1-(4-(2-pyridylmethyloxy)benzyl)-3-*i*-butyl-[(3-methyl-4-acetoxy)benzenesulfonyl]-amino-2-hydroxypropyl-carbamate (180 mg, 0.25 mmol) was dissolved in MeOH (3 mL), and K₂CO₃ (30 mg) was added. The mixture was stirred at room temperature for 30 minutes, diluted with EtOAc (50 mL), and washed with water (3x50 mL). The organic phase was

dried over MgSO_4 , concentrated, and dried *in vacuo* to yield title compound (115 mg, 70%).

^1H NMR ($\text{DMSO}-d_6$): δ 0.78 (3H, d), 0.82 (3H, d), 1.18 (1H, m), 1.27 (1H, m), 1.90 (1H, m), 2.37 (1H, t), 2.65 (3H, m), 2.91 (2H, m), 3.52 (4H, m), 3.65 (1H, t), 3.80 (1H, dd), 4.79 (1H, dd), 5.06 (2H, s), 5.46 (1H, d), 6.84 (3H, m), 7.07 (2H, d), 7.18 (1H, d), 7.30 (1H, dd), 7.38 (1H, d), 7.45 (2H, d), 7.78 (1H, d), 8.52 (1H, d), 10.31 (1H, s); MS: 670 (M^+).

Example 85

(3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[[4-(aminophenyl)sulfonyl](isobutyl)amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate (296)



To a stirred solution of (3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-(benzyloxy)benzyl]-2-hydroxy-3-{isobutyl[(4-nitrophenyl)sulfonyl]amino}propylcarbamate (0.20 g, 0.29 mmol) in 3 mL of ethanol was added 64 mg of platinum (on activated carbon, 3% Pt). The mixture was stirred under an atmospheric pressure of hydrogen for 12

hours. The catalyst was filtered, and the solvent was removed under reduced pressure. The residue was purified on silica gel using ethyl acetate-hexanes (4:1) as eluent to provide 0.10 g (52%) of the title compound.

5

¹H-NMR (DMSO-d₆): δ 0.74 (3H,d), 0.81(3H,d), 1.19 (1H,m), 1.29(1H,m), 1.89 (1H, m), 2.34 (1H,t), 2.54-2.62 (2H,m), 2.71 (1H,m), 2.90 (2H,m), 3.22 (1H,d), 3.46 (1H,m), 3.52-3.59 (3H,m), 3.66 (1H,t) 3.83 (1H,dd), 4.80 (1H,q), 4.92 (1H,d,b), 4.99 (2H,s), 5.47 (1H,d), 5.94 (2H,s), 6.54 (2H,d), 6.81 (2H,d), 7.07(2H,d), 7.19 (1H,d), 7.28 (1H,d), 7.35 (5H,m). MS: 655 (M⁺).

10

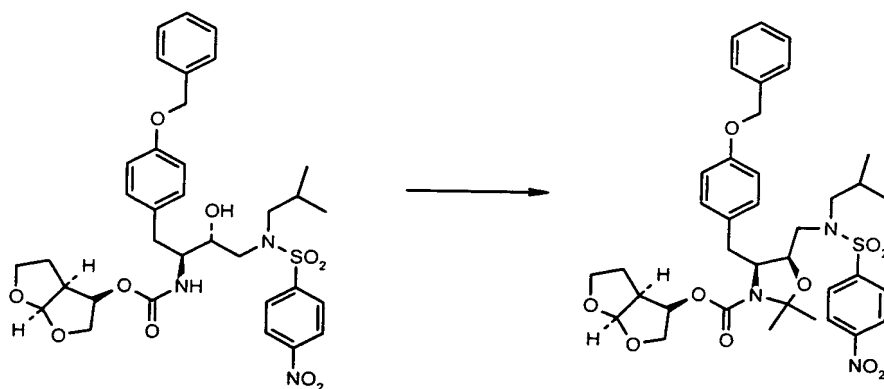
Example 86

15

Step 1:

(3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-4-[4-(benzyloxy)benzyl]-5-({isobutyl[(4-nitrophenyl)sulfonyl]amino}methyl)-2,2-dimethyl-1,3-oxazolidine-3-carboxylate.

20



To a solution of (3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-(benzyloxy)benzyl]-2-hydroxy-3-

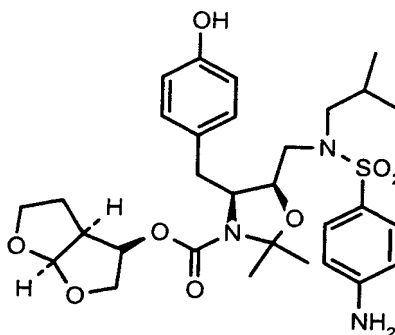
25

{isobutyl[(4-nitrophenyl)sulfonyl]amino}propylcarbamate (1.00 g, 1.5 mmol) in 20 mL of dichloromethane was added 2,2-dimethoxypropane (1.5 g, 14.6 mmol) and *p*-toluenesulfonic acid (0.09 g, 0.5 mmol). The solution was refluxed for 12 hours, and the solvent was removed under reduced pressure. The residue was purified on silica gel using ethyl acetate-hexanes (3.5:6.5) as eluent to provide 0.70 g (70%) of the title compound.

¹H-NMR (CDCl₃): δ 0.85 (3H,d), 0.90 (3H,d), 1.35,1.44 (3H,s)*, 1.55,1.61 (3H,s)*, 1.84-2.00 (3H,m,b), 2.61-2.91 (4H,m), 2.91-3.08 (3H,m), 3.17,3.33 (1H,d)*, 3.56 (1H,t), 3.80-3.86 (2H,m), 3.91-3.98 (1H,m), 4.23-4.27 (2H,m), 5.02-5.16 (2H,s), 5.67 (1H,d), 6.91 (2H,d), 7.03 (1H,d), 7.14 (1H,d), 7.31 (1H,d), 7.38 (2H,m), 7.45 (2H,d), 7.67 (2H,d), 8.25 (2H,d).

*possible indication for rotamers.

Step 2:
(3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-{[[(4-aminophenyl)sulfonyl] (isobutyl) amino]methyl}-4-(4-hydroxybenzyl)-2,2-dimethyl-1,3-oxazolidine-3-carboxylate.



To a stirred solution of (3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-
b]furan-3-yl (4*S*,5*R*)-4-[4-(benzyloxy)benzyl]-5-
({isobutyl[(4-nitrophenyl)sulfonyl]amino}methyl)-2,2-
dimethyl-1,3-oxazolidine-3-carboxylate. (0.18 g, 0.25
5 mmol) in 6 mL of a 2M solution of ammonia in methanol was
added 0.18 g of palladium (on charcoal, 10% Pd, Degussa
type). The mixture was stirred under an atmospheric
pressure of hydrogen for 12 hours. The catalyst was
10 filtered, and the solvent was removed under reduced
pressure. The residue was purified on silica gel using
ethyl acetate-hexanes (7:3) as eluent to provide 81 mg
(54%) of the title compound.

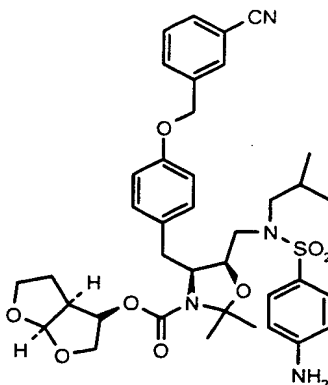
¹H-NMR (DMSO-*d*₆): δ 0.76 (3H,d), 0.80 (3H,d), 1.19,1.33
15 (3H,s)*, 1.44, 1.50 (3H,s)*, 1.74 (2H,m), 1.89 (1H,m),
2.49-2.75 (6H,m), 2.89-2.99 (2H,m), 3.19 (1H,m), 3.61
(2H,m), 3.80 (1H,m), 4.07 (1H,m), 4.74 (1H,q), 5.50,5.54
(1H,d)*, 5.94 (2H,s,b), 6.54 (2H,d), 6.64 (2H,d),
6.93,6.99 (2H,d)*, 7.13,7.20 (2H,d)*, 9.20, 9.22 (1H,s)*.
20 MS: 605 (M⁺).

*possible indication for rotamers.

Step 3:

25 (3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[[(4-aminophenyl)sulfonyl](isobutyl)amino]methyl}-4-{4-
[(3-cyanobenzyl)oxy]benzyl}-2,2-dimethyl-1,3-oxazolidine-
3-carboxylate.

30



To a stirred mixture of (3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-
 b]furan-3-yl (4*S*,5*R*)-5-{[[4-
 5 aminophenyl)sulfonyl](isobutyl)amino]methyl}-4-(4-
 hydroxybenzyl)-2,2-dimethyl-1,3-oxazolidine-3-carboxylate
 (40 mg, 0.07 mmol) and cesium carbonate (22 mg, 0.07
 mmol) in 1.5 mL of *N,N*-dimethylformamide was added 3-
 bromomethylbenzonitrile (13 mg, 0.07 mmol). The mixture
 10 was stirred at room temperature for 12 hours followed by
 dilution with 25 mL of ether. The ether was extracted
 with water (5 x 15 mL). The organic phase was dried with
 magnesium sulfate, filtered, and concentrated under
 reduced pressure. The residue was purified on silica gel
 15 using ethyl acetate-hexanes (8.5:1.5) as eluent to
 provide 41 mg (85%) of the title compound.

¹H-NMR (DMSO-*d*₆): δ 0.80-0.86 (6H,dd), 1.37 (3H,s),
 1.48,1.55 (3H,s)*, 1.75 (2H,m), 1.94 (1H,m), 2.58-2.78
 20 (6H,m), 2.92-3.09 (2H,m), 3.15-3.21 (1H,m), 3.61 (2H,m),
 3.81 (1H,m), 4.14 (1H,m), 4.74 (1H,q), 5.17 (2H,s),
 5.53,5.57 (1H,d)*, 5.99 (2H,s,b), 6.58 (2H,d), 6.95
 (2H,d), 7.11,7.14 (2H,d)*, 7.22,7.28 (2H,d)*, 7.60
 (1H,t), 7.80 (2H,m,b), 7.91 (1H,s).

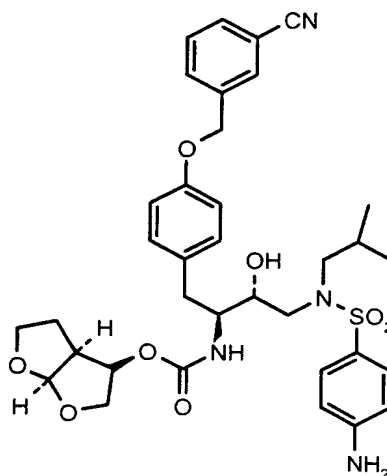
*possible indication for rotamers.

Step 4:

5

(3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[[(4-aminophenyl)sulfonyl] (isobutyl) amino]-1-{4-[(3-
cyanobenzyl)oxy]benzyl}-2-hydroxypropylcarbamate (301)

10



To a stirred solution of (3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-
b]furan-3-yl (4*S*,5*R*)-5-[[(4-
15 aminophenyl)sulfonyl] (isobutyl) amino]methyl}-4-{4-[(3-
cyanobenzyl)oxy]benzyl}-2,2-dimethyl-1,3-oxazolidine-3-
carboxylate in 3 mL of dioxane was added 4M hydrochloric
acid in dioxane (1.5 mL) and water (0.05 mL). The
solution was stirred for 1 hour followed by
20 neutralization with 10% aqueous Na₂CO₃. The solution was
extracted with dichloromethane (3 x 15 mL). The organic
phase was dried with magnesium sulfate, filtered, and
concentrated under reduced pressure. The residue was

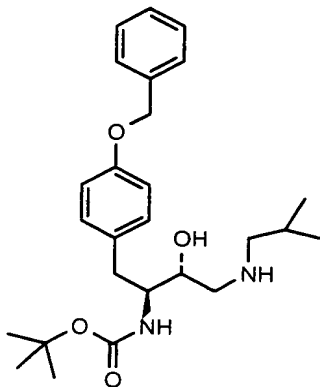
purified on silica gel using ethyl acetate-hexanes (8:2) as eluent to provide 17 mg (61%) of the title compound.

¹H-NMR (DMSO-*d*₆): δ 0.74 (3H,d), 0.81(3H,d), 1.18
5 (1H,m), 1.28 (1H,m), 1.89 (1H,m), 2.35 (1H,t), 2.54-2.60
(2H,m), 2.71 (1H,m), 2.86-2.93 (2H,m), 3.22-3.29 (1H,dd),
3.42-3.48 (1H,m), 3.53-3.59 (3H,m), 3.65 (1H,t), 3.82
(1H,q), 4.80 (1H,q), 4.93 (1H,d,b), 5.07 (2H,s), 5.47
(1H,d), 5.94 (2H,s,b), 6.54 (2H,d), 6.83 (2H,d), 7.08
10 (2H,d), 7.19 (1H,d), 7.33 (2H,d), 7.56 (1H,t), 7.74
(2H,t), 7.85 (1H,s). MS: 680 (M⁺).

Example 81

15 Step 1:

tert-butyl (1*S*,2*R*)-1-[4-(benzyloxy)benzyl]-2-hydroxy-3-(isobutylamino)propylcarbamate.



20

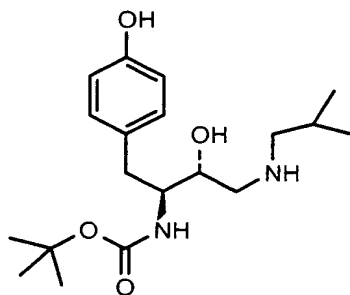
To a solution of *t*-Butyl-(1*S*,2*R*)-1-(4-benzyloxy-benzyl)-2,3-epoxydo-propyl-carbamate (3.2 g, 8.7 mmol) in 50 mL of *i*-propanol was added *i*-butylamine (4.6 g, 62.4 mmol). The mixture was stirred at 85°C for 2 hours, and then it

was cooled to 5°C followed by dropwise addition into 250 mL of water. The resulting suspension was stirred for 30 minutes at 5°C and then filtered. The solid was washed with water (3 x 150 mL) and dried under reduced pressure to yield 2.4 g (61%) of the title compound.

¹H-NMR (CDCl₃): δ 0.91 (6H,d), 1.34 (9H,s), 1.77 (1H,m), 2.37-2.48 (2H,m), 2.66-2.75 (2H,m), 2.82-2.93 (2H,m), 3.42 (1H,m), 3.77 (1H,m), 4.64 (1H,d), 5.01 (2H,s), 6.89 (2H,d), 7.12 (2H,d), 7.27-7.31 (1H,m), 7.34-7.41 (4H,m).

Step 2:

tert-butyl (1*S*,2*R*)-2-hydroxy-1-(4-hydroxybenzyl)-3-(isobutylamino)propylcarbamate.

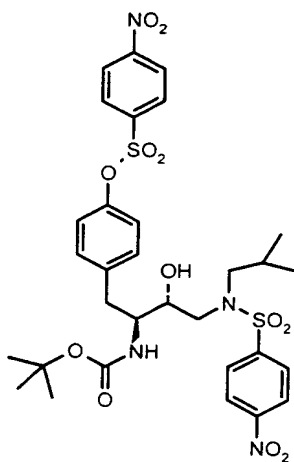


To a stirred solution of *tert*-butyl (1*S*,2*R*)-1-[4-(benzyloxy)benzyl]-2-hydroxy-3-(isobutylamino)propylcarbamate (0.5 g, 1.1 mmol) in anhydrous tetrahydrofuran (12 mL) was added 0.5 g of palladium (on charcoal, 10% Pd, Degussa type). The mixture was stirred under an atmospheric pressure of hydrogen for 12 hours. The catalyst was filtered and the solvent was removed under reduced pressure resulting in 0.4 g (98%) of the title compound.

¹H-NMR (CDCl₃): δ 0.92 (6H,dd), 1.35 (9H,s), 1.83
(1H,m), 2.44-2.55 (2H,m), 2.73-2.85 (4H,m), 3.49 (1H,m),
3.70-3.79 (1H,m,b), 4.66 (1H,d), 6.72 (2H,d), 7.05
5 (2H,d).

Step 3:

4-((2*S*,3*R*)-2-[(*tert*-butoxycarbonyl)amino]-3-hydroxy-4-
10 {isobutyl[(4-nitrophenyl)sulfonyl]-amino}butyl)phenyl 4-
nitrobenzenesulfonate



15

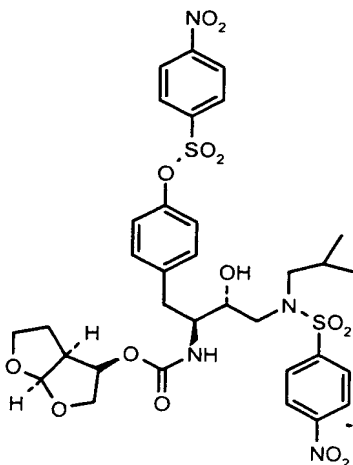
To a solution of *tert*-butyl (1*S*,2*R*)-2-hydroxy-1-(4-
hydroxybenzyl)-3-(isobutylamino)propylcarbamate (50 mg,
0.14 mmol) in 3 mL of anhydrous dichloromethane were
20 added 4-nitrobenzenesulfonyl chloride (38 mg, 0.17 mmol)
and *N*-ethyldiisopropylamine (74 mg, 0.57 mmol). After 2
hours, the solvent was removed under reduced pressure.
The residue was purified on silica gel using ethyl

acetate-hexanes (2:3) as eluent to provide 66 mg (87%) of the title compound.

¹H-NMR (DMSO-*d*₆): δ 0.77, 0.79 (6H, dd), 1.17 (9H, s),
5 1.91 (1H, m), 2.36-2.43 (1H, m), 2.84-2.95 (2H, m), 2.97-
3.01 (2H, m), 3.07-3.12 (2H, m), 3.39-3.47 (1H, m), 4.95
(1H, d), 6.67 (1H, d), 6.91 (2H, d), 7.13 (2H, d), 8.00
(2H, d), 8.07 (2H, d), 8.32 (2H, d), 8.38 (2H, d).

10 Step 4:

4-((2*S*,3*R*)-2-({[(3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-
yloxy]carbonyl}amino)-3-hydroxy-4-{isobutyl[(4-
nitrophenyl)sulfonyl]amino}butyl)phenyl 4-
15 nitrobenzenesulfonate (297)



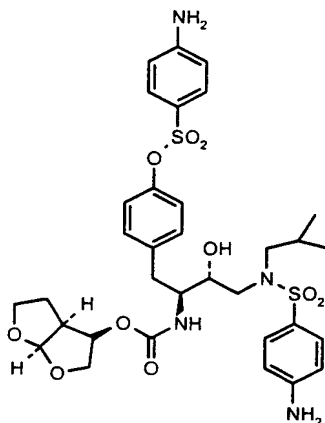
To a solution of 4-((2*S*,3*R*)-2-[(*tert*-
20 butoxycarbonyl)amino]-3-hydroxy-4-{isobutyl[(4-
nitrophenyl)sulfonyl]-amino}butyl)phenyl 4-
nitrobenzenesulfonate (0.21 mg, 0.29 mmol) in 10 mL of
dichloromethane was added dropwise trifluoroacetic acid

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(2 mL). The mixture was stirred for 1 hour at room temperature. The solvents were removed under reduced pressure; the residue was redissolved in 50 mL of dichloromethane and 20 mL of 5% aqueous sodium carbonate was added. The aqueous phase was separated and extracted with dichloromethane (2 x 20 mL). The combined organic phases were dried over sodium carbonate, filtered, and concentrated under reduced pressure. The residue was redissolved in 10 mL of acetonitrile. N-Ethyldiisopropylamine (80 uL, 0.05 mmol) and (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl-4-nitrophenylcarbonate (0.11g, 0.37 mmol) were added, and the solution was stirred for 12 hours. The solvents were removed under reduced pressure. The residue was redissolved in ethyl acetate (30 mL), and the organic phase was extracted with water (20 mL) followed by 5% aqueous sodium carbonate solution (5 x 20 mL). The organic phase was dried with sodium carbonate, filtered, and concentrated under reduced pressure. The residue was purified on silica gel using ethyl acetate-hexanes as eluent to provide 0.12 g (55%) of the title compound.

¹H-NMR (CD₃OD): δ 0.84 (3H,d), 0.90 (3H,d), 1.46 (1H,m), 1.58 (1H,m), 1.99 (1H,m), 2.47 (1H,t), 2.87-2.97 (2H,m), 3.04-3.10 (2H,m), 3.18 (1H,m), 3.44-3.47 (1H,dd), 3.62-3.68 (4H,m), 3.79 (1H,m), 3.89 (1H,q), 4.91 (1H,q), 5.57 (1H,d), 6.91 (2H,d), 7.20 (2H,d), 8.02-8.05 (4H,d+d), 8.36-8.42 (4H,d+d). MS: 778 (M⁺).

(3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(4-aminobenzyl)(isobutyl)amino]-1-{4-[(4-
 aminobenzyl)oxy]benzyl}-2-hydroxypropylcarbamate (298)



5

To a stirred solution of 4-((2*S*,3*R*)-2-(([(3*S*,3*aS*,6*aR*)-
 hexahydrofuro[2,3-*b*]furan-3-yloxy]carbonyl)amino)-3-
 hydroxy-4-{isobutyl[(4-
 10 nitrophenyl)sulfonyl]amino}butyl)phenyl 4-
 nitrobenzenesulfonate (0.12 g, 0.15 mmol) in 2 mL of
 anhydrous tetrahydrofuran was added 36 mg of platinum (on
 activated carbon, 3% Pt). The mixture was stirred under
 an atmospheric pressure of hydrogen for 3 hours. The
 15 catalyst was filtered, and the solvent was removed under
 reduced pressure. The residue was purified on silica gel
 using ethyl acetate-hexanes (9:1) as eluent to provide 39
 mg (35%) of the title compound.

20 ¹H-NMR (CD₃OD): δ 0.84 (3H,d), 0.90 (3H,d), 1.38-1.43
 (1H,m), 1.53-1.59 (1H,m), 1.96 (1H,m), 2.47 (1H,t), 2.70-
 2.88 (4H,m), 2.98 (1H,m), 3.13,3.21 (1H,dd), 3.31,3.35
 (1H,dd), 3.64-3.71 (4H,m), 3.73-3.82 (3H,m), 3.85-3.92
 (1H,m), 4.91 (1H,q), 5.57 (1H,d), 6.62,6.66 (4H,d+d),

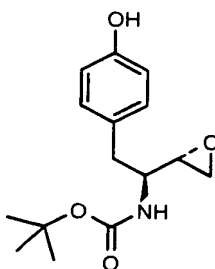
6.82 (2H,d), 7.16 (2H,d), 7.38 (2H,d), 7.45 (2H,d). MS:
720 (M⁺).

5

Example 83

Step 1:

tert-butyl (1S)-2-(4-hydroxyphenyl)-1-[(2S)-
oxirany]ethylcarbamate.



10

To a stirred solution of t-Butyl-(1S,2R)-1-(4-benzyloxy-
benzyl)2,3-epoxydo-propylcarbamate in a mixture of
ethanol-ethyl acetate (4:1) was added palladium hydroxide
15 (on carbon, 20% palladium). The mixture was stirred under
an atmospheric pressure of hydrogen for 2 hours. The
catalyst was filtered and washed one time each with
ethanol, methanol, and ethyl acetate. The solvent was
removed under reduced pressure providing 0.5 g
20 (quantitative) of the title compound.

25

¹H-NMR (DMSO-d₆): δ 1.30 (9H,s), 2.59-2.75 (4H,m), 2.85
(1H,m), 3.35-3.46 (1H,m), 6.63 (2H,d), 6.78 (1H,d), 6.98
(2H,d), 9.03,9.12 (1H,s)*.

*possible indication for rotamers.

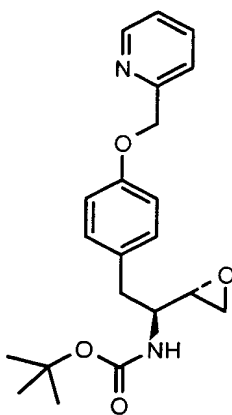
Step 2:

General procedure for the alkylation of product from Procedure 11.

5

To a stirred mixture of *tert*-butyl (1*S*)-2-(4-hydroxyphenyl)-1-[(2*S*)-oxiranyl]ethylcarbamate (0.25 g, 0.90 mmol) and cesium carbonate (0.73 g, 2.2 mmol) in 5 mL of *N,N*-dimethylformamide was added the desired alkyl
10 halide [1] (0.90 mmol). The mixture was stirred at room temperature for 12 hours followed by dilution with 125 mL of ether. The ether was extracted with water (5 x 75 mL). The organic phase was dried with magnesium sulfate, filtered, and concentrated under reduced pressure. The
15 residue was purified on silica gel using ethyl acetate-hexanes [2] as eluent to provide the following compounds [Y:(3)]:

1. *tert*-butyl (1*S*)-1-[(2*S*)-oxiranyl]-2-[4-(2-pyridinylmethoxy)phenyl]ethylcarbamate.
20



[1]: 2-Picolyl chloride hydrochloride; [2]: no purification; [3]: 91%.

$^1\text{H-NMR}$ ($\text{DMSO}-d_6$): δ 1.23 (9H,s), 2.56-2.63 (3H,m), 2.71-2.76 (1H,m), 2.85 (1H,m), 3.40 (1H,m), 5.08 (2H,s), 6.79 (1H,d), 6.87 (2H,d), 7.07 (2H,d), 7.29 (1H,m), 7.44 (1H,d), 7.77 (1H,t), 8.52 (1H,d).

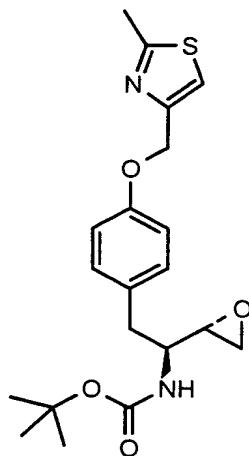
5

Example 84

Step 1

2. *tert*-butyl (1*S*)-2-{4-[(2-methyl-1,3-thiazol-4-yl)methoxy]phenyl}-1-[(2*S*)-oxiranyl]-ethylcarbamate.

10



[1]: 4-Chloromethyl-2-methylthiazole hydrochloride*;

15 [2]: (6:4); [3]: 35%.

$^1\text{H-NMR}$ ($\text{DMSO}-d_6$): δ 1.24 (9H,s), 2.57-2.63 (6H,m+s), 2.72, 2.76 (1H, dd), 2.84 (1H,m), 3.41 (1H,m), 5.00 (2H,s), 6.79 (1H,d), 6.87 (2H,d), 7.07 (2H,d), 7.46 (1H,s).

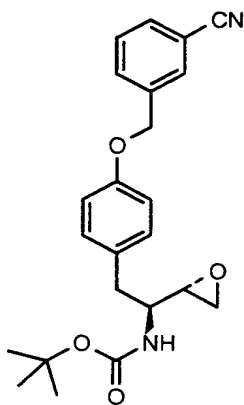
20

*added excess sodium iodide to the reaction.

Example 85

5 Step 1:

3. *tert*-butyl (1*S*)-2-{4-[(3-cyanobenzyl)oxy]phenyl}-1-[(2*S*)-oxiranyl]ethylcarbamate.



[1]: 3-(Bromomethyl)benzonitrile; [2]: (1:1); [3]:
75%.

15 ¹H-NMR (DMSO-*d*₆): δ 1.23 (9H,s), 2.55-2.63 (3H,m),
2.72,2.75 (1H,dd), 2.84 (1H,m), 3.41 (1H,m), 5.09 (2H,s),
6.79 (1H,d), 6.88 (2H,d), 7.08 (2H,d), 7.56 (1H,t), 7.74
(2H,t), 7.84 (1H,s).

20

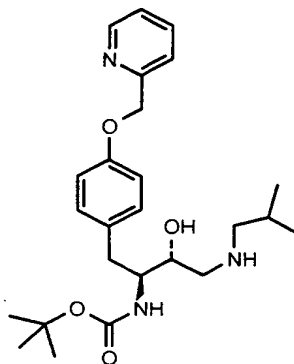
Example 83, Step 3:

General procedure for the ring-opening

To a solution of the product from the previous step (0.4
25 g, 1.2 mmol) in *i*-propanol was added *i*-butylamine (0.4 g,

5.8 mmol). The mixture was stirred at 85°C for 2 hours, and then it was cooled to 5°C followed by dropwise addition into 50 mL of water. The resulting suspension was stirred for 30 minutes at 5°C and then filtered. The solid was washed with water (3 x 25 mL) and dried under reduced pressure to provide the following compounds [Y:(1)]:

1. *tert*-butyl (1*S*,2*R*)-2-hydroxy-3-(isobutylamino)-1-[4-(2-pyridinylmethoxy)benzyl]-propylcarbamate.

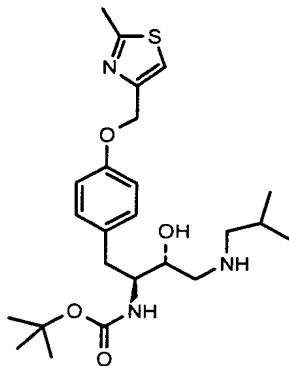


[1]: 82%.

¹H-NMR (DMSO-*d*₆): δ 0.82 (6H,dd), 1.21 (9H,s), 1.59 (1H,m), 2.25 (2H,d), 2.39-2.42 (2H,m), 2.51 (1H,m), 2.85 (1H,dd), 3.34-3.43 (3H,m), 4.71 (1H,s,b), 5.08 (2H,s), 6.65 (1H,d), 6.85 (2H,d), 7.04 (2H,d), 7.29 (1H,m), 7.43 (1H,d), 7.77 (1H,t), 8.52 (1H,d).

Example 84, Step 2:

tert-butyl (1*S*,2*R*)-2-hydroxy-3-(isobutylamino)-1-{4-[(2-methyl-1,3-thiazol-4-yl)methoxy]benzyl}propylcarbamate.



5

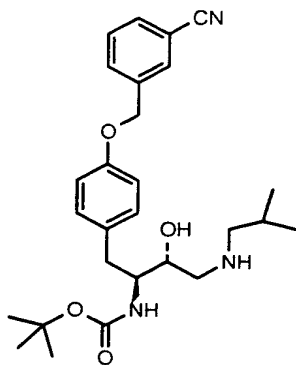
[1]: 66%.

¹H-NMR (DMSO-*d*₆): δ 0.81 (6H,d), 1.22 (9H,s), 1.60 (1H,m), 2.27 (2H,m), 2.40-2.55 (2H,m,b), 2.60 (3H,s),
 10 2.86 (1H,d,b), 3.35-3.42 (3H,m), 4.73 (1H,s,b), 4.99 (2H,s), 6.65 (1H,d), 6.85 (2H,d), 7.03 (2H,d), 7.45 (1H,s).

Example 85, Step 2:

15

tert-butyl (1*S*,2*R*)-1-{4-[(3-cyanobenzyl)oxy]benzyl}-2-hydroxy-3-(isobutylamino)-propylcarbamate.



[1]: 88%.

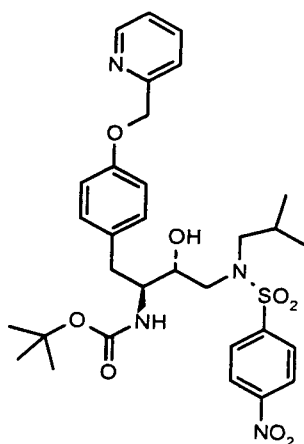
¹H-NMR (DMSO-d₆): δ 0.80 (6H,d), 1.21 (9H,s), 1.59
5 (1H,m), 2.25 (2H,d), 2.43 (2H,m), 2.51 (1H,m), 2.86
(1H,dd), 3.34 (1H,m), 3.41 (1H,m), 4.70 (1H,s,b), 5.07
(2H,s), 6.64 (1H,d), 6.85 (2H,d), 7.04 (2H,d), 7.55
(1H,t), 7.74 (2H,t), 7.84 (1H,s).

10 **Example 83, Step 4:**

General procedure for the sulfonylation

To a stirred solution of the product from the previous
step (0.42 g, 0.96 mmol) in 20 mL of anhydrous
15 dichloromethane were added 4-nitrobenzenesulfonyl
chloride (0.25 g, 1.1 mmol) and N-ethyldiisopropylamine
(0.15 g, 1.7 mmol). The mixture was allowed to react for
the given number of hours [1] and at which time the
solvent was removed under reduced pressure. The residue
20 was redissolved in dichloromethane and extracted with
water (3 x 50 mL). The organic phase was dried with
magnesium sulfate, filtered, and removed under reduced
pressure. The residue was purified on silica gel using
ethyl acetate-hexanes [2] as eluent to provide the
25 following compounds [Y:(3)]:

1. tert-butyl (1S,2R)-2-hydroxy-3-{isobutyl[(4-
nitrophenyl)sulfonyl]amino}-1-[4-(2-
pyridinylmethoxy)benzyl]propylcarbamate.

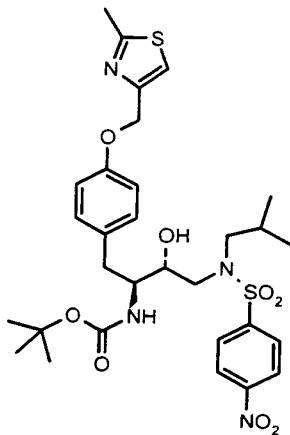


[1]: 6 hours; [2]: (7:3); [3]: 87%.

¹H-NMR (DMSO-*d*₆): δ 0.77, 0.80 (6H, dd), 1.20 (9H, s), 1.92 (1H, m), 2.36 (2H, d), 2.80-2.91 (2H, m), 2.98-3.13 (2H, m), 3.42-3.46 (2H, m), 4.89 (1H, d), 5.07 (2H, s), 6.61 (1H, d), 6.84 (2H, d), 7.03 (2H, d), 7.29 (1H, m), 7.43 (1H, d), 7.76 (1H, m), 8.00 (2H, d), 8.32 (2H, d), 8.52 (1H, d).

Example 84, Step 3:

tert-butyl (1S,2R)-2-hydroxy-3-{isobutyl[(4-nitrophenyl)sulfonyl]amino}-1-{4-[(2-methyl-1,3-thiazol-4-yl)methoxy]benzyl}propylcarbamate.

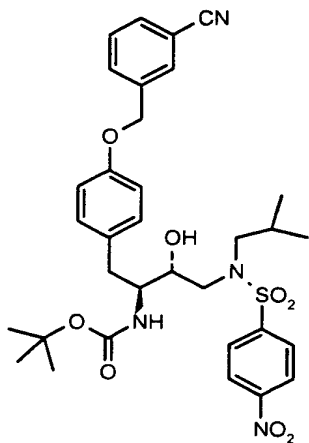


[1]: 12 hours; [2]: (1:1); [3]: 83%.

¹H-NMR (DMSO-d₆): δ 0.79 (6H,dd), 1.21 (9H,s), 1.93
5 (1H,m), 2.34-2.40 (2H,m), 2.60 (3H,s), 2.80-2.91 (2H,m),
2.99-3.13 (2H,m), 3.33 (1H,m), 3.44 (1H,m), 4.89 (1H,d),
4.99 (2H,s), 6.63 (1H,d), 6.85 (2H,d), 7.04 (2H,d), 7.45
(1H,s), 8.01 (2H,d), 8.33 (2H,d).

10 **Example 85, Step 3:**

**tert-butyl (1S,2R)-1-{4-[(3-cyanobenzyl)oxy]benzyl}-2-
hydroxy-3-{isobutyl[(4-
15 nitrophenyl)sulfonyl]amino}propylcarbamate.**



[1]: 12 hours; [2]: (1:1); [3]: 95%.

20 ¹H-NMR (DMSO-d₆): δ 0.78 (6H,d+d), 1.19 (9H,s), 1.92
(1H,m), 2.36 (2H,m), 2.80-2.91 (2H,m), 2.98-3.13 (2H,m),
3.32 (1H,m), 3.42 (1H,m), 4.89 (1H,d), 5.07 (2H,s), 6.61
(1H,d), 6.84 (2H,d), 7.04 (2H,d), 7.55 (1H,t), 7.74
(2H,t), 7.83 (1H,s), 8.00 (2H,d), 8.31 (2H,d).

Example 83, Step 5

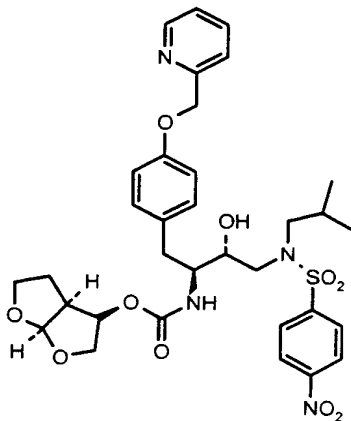
5

General procedure for the addition of BisTHF units

006090 1946560

To a solution of the product from the previous step (0.36 g, 0.57 mmol) in 20 mL of anhydrous dichloromethane was added dropwise trifluoroacetic acid (5 mL). The mixture was stirred for 1 hour at room temperature. The solvents were removed under reduced pressure; the residue was redissolved in 50 mL of dichloromethane and 30 mL of 10% aqueous sodium carbonate was added. The aqueous phase was separated and extracted with dichloromethane (2 x 25 mL). The combined organic phases were dried with sodium carbonate, filtered, and concentrated under reduced pressure. The residue was redissolved in 10 mL of acetonitrile. N-Ethyldiisopropylamine (0.15 g, 1.1 mmol) and (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl-4-nitrophenylcarbonate (0.20 g, 0.68 mmol) were added, and the solution was stirred for 12 hours. The solvents were removed under reduced pressure. The residue was redissolved in ethyl acetate (50mL), and the organic phase was extracted with water (30 mL) followed by 5% aqueous sodium carbonate solution (5 x 50 mL). The organic phase was dried with sodium carbonate, filtered, and concentrated under reduced pressure. The residue was purified on silica gel using ethyl acetate-hexanes [1] as eluent to provide the following compounds [Y:(2)]:

1. (3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-2-hydroxy-3-{isobutyl[(4-nitrophenyl)sulfonyl]amino}-1-[4-(2-pyridinylmethoxy)benzyl]propylcarbamate.



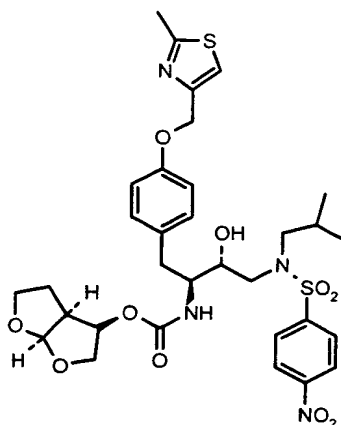
5

[1]: (9:1→100%); [2]: Y: 77%.

¹H-NMR (DMSO-*d*₆): δ 0.77 (3H,d), 0.81 (3H,d), 1.20 (1H,m), 1.32 (1H,m), 1.92 (1H,m), 2.32 (1H,t), 2.72 (1H,m), 2.84-2.89 (2H,dd,b), 2.95-3.00 (1H,m), 3.08-3.14 (1H,m), 3.33 (1H,d), 3.45 (2H,m,b), 3.50-3.57 (2H,m), 3.65 (1H,t), 3.80 (1H,q), 4.80 (1H,q), 4.98 (1H,d), 5.06 (2H,s), 5.46 (1H,d), 6.83 (2H,d), 7.05 (2H,d), 7.18 (1H,d), 7.29 (1H,m), 7.44 (1H,d), 7.78 (1H,m), 8.01 (2H,d), 8.34 (2H,d), 8.52 (1H,d).

Example 84, Step 4:

(3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-2-hydroxy-3-{isobutyl[(4-nitrophenyl)sulfonyl]amino}-1-{4-[(2-methyl-1,3-thiazol-4-yl)methoxy]benzyl}propylcarbamate.

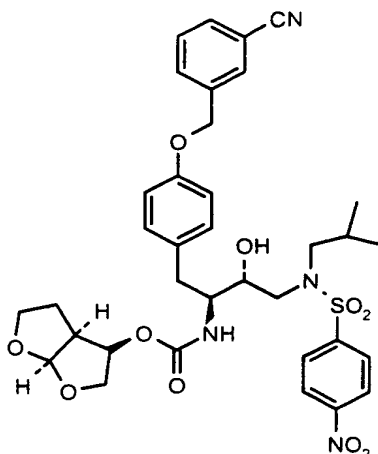


[1]: (8:2); [2]: 59%.

5 $^1\text{H-NMR}$ ($\text{DMSO}-d_6$): δ 0.77 (3H,d), 0.81 (3H,d), 1.20
 (1H,m), 1.29-1.36 (1H,m), 1.92 (1H,m), 2.29-2.35 (1H,m),
 2.60 (3H,s), 2.71-2.76 (1H,m), 2.85-2.89 (2H,dd), 2.95-
 3.01 (1H,m), 3.08-3.14 (1H,m), 3.33 (1H,d), 3.45
 (2H,m,b), 3.50-3.58 (2H,m), 3.67 (1H,t), 3.80 (1H,q),
 10 4.80 (1H,q), 4.98 (3H,s,b), 5.47 (1H,d), 6.83 (2H,d),
 7.05 (2H,d), 7.18 (1H,d), 7.47 (1H,s), 8.01 (2H,d), 8.34
 (2H,d).

Example 85, Step 4:

(3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-{4-
 15 [(3-cyanobenzyl)oxy]benzyl}-2-hydroxy-3-{isobutyl[(4-
 nitrophenyl)sulfonyl]amino}propylcarbamate.



[1]: (2:8); [2]: 74%.

5

¹H-NMR (DMSO-*d*₆): δ 0.76 (3H,d), 0.80 (3H,d), 1.16-1.20
 10 (1H,m), 1.27-1.37 (1H,m), 1.92 (1H,m), 2.32 (1H,t), 2.72
 (1H,m), 2.86 (2H,dd), 2.94-3.00 (1H,m), 3.08-3.14 (1H,m),
 3.33 (1H,d), 3.45 (2H,m,b), 3.50-3.58 (2H,m), 3.65
 (1H,t), 3.80 (1H,q), 4.80 (1H,q), 4.98 (1H,d), 5.06
 (2H,s), 5.46 (1H,d), 6.83 (2H,d), 7.06 (2H,d), 7.18
 15 (1H,d), 7.56 (1H,t), 7.74 (2H,t), 7.84 (1H,s), 8.00
 (2H,d), 8.33 (2H,d).

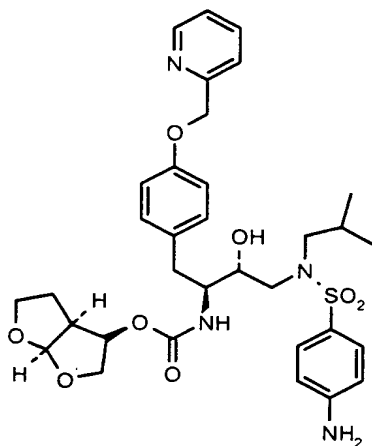
Example 83, Step 6

20 General procedure for the reduction of the nitro group

To a stirred solution of the product obtained from the previous step (0.23 g, 0.34 mmol) in 4 mL of anhydrous tetrahydrofuran was added 70 mg of platinum (on activated

carbon, 3% Pt). The mixture was stirred under an atmospheric pressure of hydrogen for the indicated number of hours [1]. The catalyst was filtered, and the solvent was removed under reduced pressure. The residue was
5 purified on silica gel using ethyl acetate-hexanes [2] as eluent to provide the following compounds (Y:[3]):

(3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
10 [[(4-aminophenyl)sulfonyl]-(isobutyl)amino]-2-hydroxy-1-
[4-(2-pyridinylmethoxy)benzyl]propylcarbamate (299)



15

[1]: 12 hours; [2]: (9:1); [3]: 90%*.

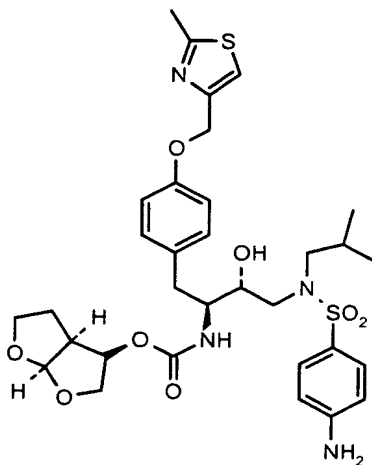
¹H-NMR (DMSO-*d*₆): δ 0.74 (3H,d), 0.81 (3H,d), 1.20-1.16
(1H,m), 1.29 (1H,m), 1.89 (1H,m), 2.34 (1H,t), 2.54-2.66
20 (2H,m), 2.71 (1H,m), 2.86-2.96 (2H,m), 3.22,3.25 (1H,dd),
3.40-3.59 (4H,m,b), 3.65 (1H,t), 3.82 (1H,m), 4.80
(1H,q), 4.92,4.95 (1H,dd), 5.06,5.08 (2H,s), 5.46 (1H,d),
5.94 (1H,s), 6.54 (1H,d), 6.82 (3H,m), 7.08 (2H,d), 7.20
(1H,m), 7.31 (2H,m), 7.44 (1H,d), 7.49 (1H,d), 7.78

(1H,t), 8.52 (2H,d), 8.64 (s)*, 8.94 (s)*. MS: 657 (M⁺), 673 (M⁺)*.

*Note: This was obtained as a 3:1 mixture with respect
5 to the hydroxylamine derivative (by ¹H NMR).

Example 84 Step 5:

(3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
10 [[(4-aminophenyl)sulfonyl]-(isobutyl)amino]-2-hydroxy-1-
{4-[(2-methyl-1,3-thiazol-4-yl)methoxy]benzyl}
propylcarbamate (300)



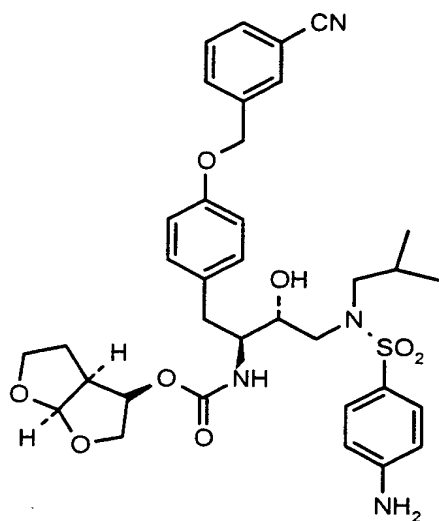
15

[1]: 77 hours; [2]: (9:1); [3]: 53%.

¹H-NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.19
(1H,m), 1.32 (1H,m), 1.89 (1H,m), 2.35 (1H,t), 2.54-2.69
20 (5H, dd+s), 2.72 (1H,m), 2.87-2.93 (2H,m), 3.22 (1H,d),
3.47 (1H,m), 3.53-3.60 (3H,m), 3.68 (1H,t), 3.82 (1H,dd)
4.80 (1H,q), 4.93 (1H,m), 4.98 (2H,s,b), 5.47 (1H,d),
5.94 (2H,s,b), 6.55 (2H,d), 6.83 (2H,d), 7.07 (2H,d),
7.19 (1H,d), 7.33 (2H,d), 7.47 (1H,s). MS: 676 (M⁺).

Example 85, Step 5:

- 5 (3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[[4-aminophenyl)sulfonyl]-(isobutyl)amino]-1-{4-[(3-
cyanobenzyl)oxy]benzyl}-2-hydroxypropylcarbamate (301)

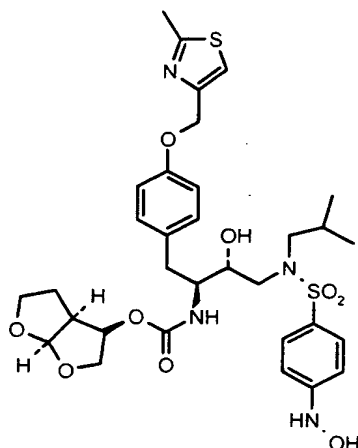


analytical data see Example 301 (Route 1) above

10

Example 86

- (3*S*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-2-
hydroxy-3-[[4-(hydroxyamino)benzyl]-(isobutyl)amino]-1-{4-
15 [(2-methyl-1,3-thiazol-4-yl)methoxy]benzyl}
propylcarbamate (302)



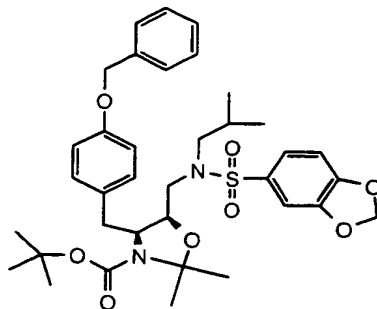
[1]: 3 hours; [2]: no purification; [3]: quantitative.

5 $^1\text{H-NMR}$ ($\text{DMSO}-d_6$): δ 0.74 (3H,d), 0.81 (3H,d), 1.15-1.20
 (1H,m), 1.34 (1H,m), 1.90 (1H,m), 2.35 (1H,t), 2.60-2.75
 (7H, m+s), 2.87-2.96 (1H,m), 3.29 (1H,d,b), 3.42-3.56
 (4H,m), 3.68 (1H,t), 3.82 (1H,dd), 4.80 (1H,q), 4.98,4.95
 (3H,s+d), 5.47 (1H,d), 6.82,6.84 (4H,d+d), 7.07 (2H,d),
 10 7.20 (1H,d), 7.48,7.51 (3H,sd), 8.64 (1H,s), 8.94 (1H,s).
 MS: 676 (M^+).

Example 87

Step 1:

15 **t-butyl (4S,5R)-5-{[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-4-[4-(benzyloxy)benzyl]-2,2-dimethyl-1,3-oxazolidine-3-carboxylate**



The reaction was carried out as described for analogous transformations above, starting with previously described
 5 tert-butyl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate

The residue was purified on silica gel hexane-ethyl acetate (3:1) as eluant to afford 2.68g (88%) of the
 10 title compound.

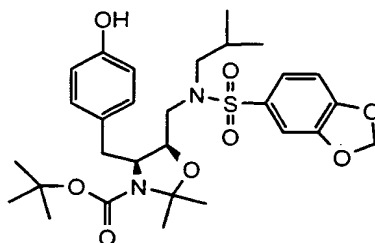
¹H-NMR: (CDCl₃): δ 0.86 (3H,d), 0.98 (3H,d), 1.40, 1.45 (3H,s)*, 1.50, 1.55 (3H,s)*, 1.64 (3H,s), 1.66 (3H,s), 1.68 (3H,s), 1.99 (1H,m), 2.65-3.09 (5H,m), 3.24-3.32
 15 (1H,m), 4.17-4.28 (2H,m), 5.09 (2H,s), 6.05 (2H,s), 6.82 (1H,d), 6.96 (2H,d), 7.05-7.18 (4H,m), 7.29-7.50 (5H,m).
 MS: 667 (M⁺). C₃₆H₄₆N₂O₈S.

*: Possible indication for rotamers.

20

Step 2:

t-butyl (4*S*,5*R*)-5-{[1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-4-(4-hydroxybenzyl)-
 25 2,2-dimethyl-1,3-oxazolidine-3-carboxylate.



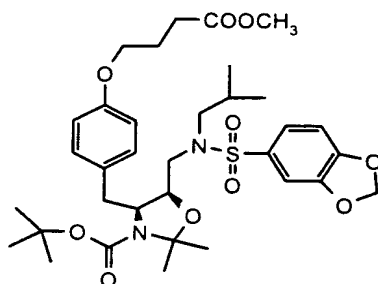
The reaction was carried out as described previously
 5 starting from t-butyl (4S,5R)-5-{[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-4-[4-(benzyloxy)benzyl]-2,2-dimethyl-1,3-oxazolidine-3-carboxylate to afford 2.31 g (100%) of the title compound.

¹H-NMR: (CDCl₃): δ 0.86 (3H,d), 0.98 (3H,d), 1.40, 1.45
 (3H,s)*, 1.50, 1.55 (3H,s)*, 1.66 (3H,s), 1.68 (3H,s),
 1.70 (3H,s), 1.99 (1H,m), 2.63-3.09 (5H,m), 3.23-3.31
 (1H,m), 3.80 (1H,m), 4.14-4.27 (2H,m), 6.11 (2H,s), 6.77-
 15 6.87 (3H,m), 7.02-7.19 (4H,m). MS: 576 (M⁺). C₂₉H₄₀N₂O₈S.

*: Possible indication for rotamers.

20 Step 3:

t-butyl (4S,5R)-5-{[1,3-benzodioxol-5-ylsulfonyl](isobutyl)amino]methyl}-4-{4-(4-methoxy-4-oxobutoxy)benzyl}-2,2-dimethyl-1,3-oxazolidine-3-
 25 carboxylate



The reaction was carried out as described previously for
 5 analogous transformations, starting from 1.83 g (3.17
 mmol) of t-butyl (4S,5R)-5-{[1,3-benzodioxol-5-
 ylsulfonyl] (isobutyl) amino] methyl}-4-(4-hydroxybenzyl)-
 2,2-dimethyl-1,3-oxazolidine-3-carboxylate except methyl-
 4-iodobutyrate was used as alkylating reagent.

10

[2]: room temperature; [3]: 5 hours.

The residue was purified on silica gel using hexane-ethyl
 15 acetate (2:1/1:1) as eluant to afford 2.1g (98%) of the
 title compound.

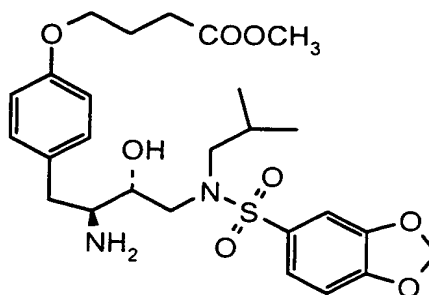
¹H-NMR: (CDCl₃): δ 0.86 (3H,d), 0.98 (3H,d), 1.36, 1.38
 (3H,s)*, 1.44, 1.49 (3H,s)*, 1.56 (3H,s), 1.59 (3H,s),
 1.64 (3H,s), 1.99 (1H,m) 2.14-2.19 (1H,m), 2.47-3.08
 20 (7H,m), 3.23-3.30 (2H,m), 3.72 (3H,s), 4.03-4.27 (4H,m),
 6.10 (2H,s), 6.80-6.87 (3H,m), 7.06-7.19 (4H,m). MS: 677
 (M⁺). C₃₄H₄₈N₂O₁₀S.

*: Possible indication for rotamers.

25

Step 4:

Methyl 4-(4-{(2S,3R)-2-amino-4-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenoxy)butanoate



5

The reaction was carried out as described previously for similar transformations starting t-butyl (4S,5R)-5-{[1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-4-{4-(4-methoxy-4-oxobutoxy)benzyl}-2,2-dimethyl-1,3-oxazolidine-3-carboxylate, to afford 1.6g (96%) of the title compound.

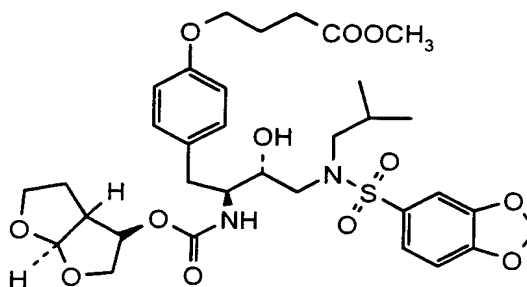
Used in the next step without further purification.

15 ¹H-NMR: (Methanol-d₄): δ 0.86 (3H,d), 0.98 (3H,d), 1.99 (1H,m), 2.02-2.92 (3H,m), 2.45-2.60 (2H,m), 2.67-3.17 (6H,m), 3.28-3.47 (2H,m), 3.57-3.62 (1H,m), 3.70 (3H,s), 3.75-3.82 (1H,m), 3.92-4.18 (2H,m), 6.14 (2H,s), 6.81-7.03 (4H,m), 7.13-7.22 (2H,m), 7.39 (1H,d). MS: 537 (M⁺).
20 C₂₆H₃₆N₂O₈S.

Step 5:

Methyl 4-(4-{(2S,3R)-2-({[3R,3aS,6aR]-hexahydrofuro[2,3-b]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-

ylsulfonyl) (isobutyl) amino] -3-hydroxybutyl}phenoxy)
butanoate (303)



5

The reaction was carried out as described previously for
analogous transformations, starting Methyl 4-(4-{(2S,3R)-
2-amino-4-[(1,3-benzodioxol-5-
10 ylsulfonyl) (isobutyl) amino] -3-
hydroxybutyl}phenoxy)butanoate

The residue was purified by silica gel using hexane-ethyl
acetate (2:1/1:1/1:2) as eluant to afford 0.91 g (44%) of
the title compound.

15

¹H-NMR: (CDCl₃): δ 0.94 (3H,d), 1.00 (3H,d), 1.25-1.35
(1H,m), 1.56-1.69 (2H,m), 1.89 (1H,m), 2.08-2.18 (3H,m),
2.53-2.58 (2H,m), 2.75-2.82 (2H,m), 2.84-3.13 (4H,m),
3.18-3.21 (2H,m), 3.61 (1H,s), 3.72 (3H,s), 3.78-3.97
20 (2H,m), 4.01-4.17 (3H,m), 4.90 (1H,d), 5.06 (1H,q), 5.70
(1H,d), 6.13 (2H,s), 6.83 (2H,d), 6.93 (1H,d), 7.13-7.19
(3H,m), 7.37 (1H,d). MS: 693 (M⁺). C₃₃H₄₄N₂O₁₂S.

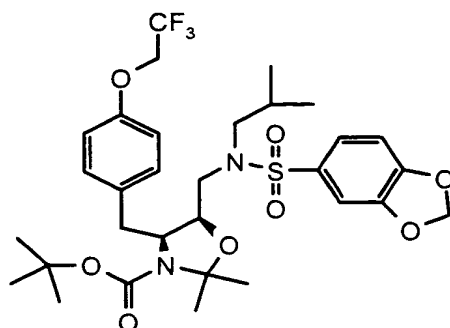
25

Example 88

Step 1:

t-butyl (4S,5R)-5-{[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-2,2-dimethyl-4-[4-(2,2,2-trifluoroethoxy)benzyl]-1,3-oxazolidine-3-carboxylate

5



To a solution of t-butyl (4S,5R)-5-{[1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-4-(4-hydroxybenzyl)-2,2-dimethyl-1,3-oxazolidine-3-carboxylate (0.25 g, 0.433 mmol) in 6 ml of dichloromethane-THF (2:1) was added tetrabutylammonium hydrogensulfate (8.9 mg, 0.026 mmol), 40% aqueous sodium hydroxide (0.136 ml) and 2,2,2-trifluroethyltrifluoromethanesulfonate (100 mg, 0.433 mmol). The mixture was heated to reflux for 2.5 hours. Diluted with 10 ml of dichloromethane and 10 ml of water and the organic phase was separated, dried with sodium sulfate, filtered and concentrated under reduced pressure.

The residue was purified on silica gel using hexane-ethyl acetate (2:1) as eluant to afford 90 mg (32%) of the title compound.

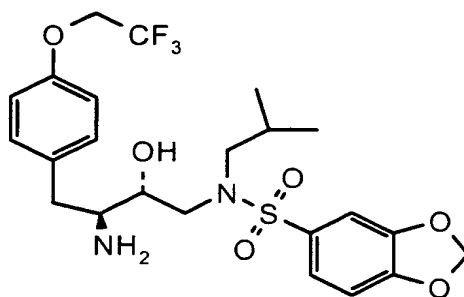
¹H-NMR: (CDCl₃): δ 0.87 (3H,d), 0.96 (3H,d), 1.45, 1.48 (3H,s)*, 1.51,1.57 (3H,s)*, 1.60 (3H,s), 1.62(3H,s), 1.65

(3H,s), 1.99 (1H,m), 2.68-3.19 (5H,m), 3.25-3.30 (1H,m),
4.26-4.40 ((4H,m), 6.09 (2H,s), 6.81-7.01 (4H,m), 7.17-
7.29 (3H,m). MS: 659 (M⁺). C₃₁H₄₁F₃N₂O₈S.

5 *: Possible indication for rotamers.

Step 2:

N-{(2R,3S)-3-amino-2-hydroxy-4-[4-(2,2,2-
10 trifluoroethoxy)phenyl]butyl}-N-isobutyl-1,3-
benzodioxole-5-sulfonamide



15

The reaction was carried out as previously for similar
transformations from t-butyl (4S,5R)-5-{[(1,3-
benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-2,2-
dimethyl-4-[4-(2,2,2-trifluoroethoxy)benzyl]-1,3-
20 oxazolidine-3-carboxylate to afford 70 mg (99%) of the
title compound.

Used in the next step without further purification.

¹H-NMR: (DMSO-d₆): δ 0.78 (3H,d), 0.84 (3H,d), 1.99
25 (1H,m), 2.60-3.58 (11H,m), 4.72-4.84 (2H,m), 6.16 (2H,s),

6.93-7.18 (4H,m), 7.21-7.34 (3H,m). MS: 519 (M⁺).

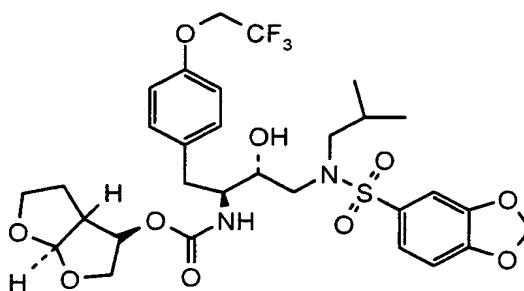
C₂₃H₂₉F₃N₂O₆S.

Step 3:

5

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-[4-(2,2,2-trifluoroethoxy)benzyl]
propylcarbamate (304)

10



15 The reaction was carried out as described previously for
similar transformations from N-{(2R,3S)-3-amino-2-
hydroxy-4-[4-(2,2,2-trifluoroethoxy)phenyl]butyl}-N-
isobutyl-1,3-benzodioxole-5-sulfonamide
The residue was purified by silica gel using hexane-ethyl
20 acetate (1:1) as eluant to afford 43 mg (47%) of the
title compound.

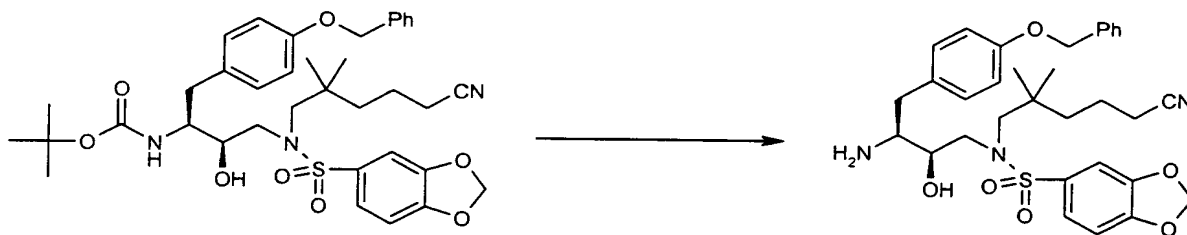
¹H-NMR: (CDCl₃): δ 0.86 (3H,d), 0.95 (3H,d), 1.31-1.39
(1H,m), 1.59-1.88 (4H,m), 2.79-2.86 (2H,m), 2.97-3.08
25 (2H,m), 3.12-3.20 (2H,m), 3.64-4.06 (6H,m), 4.32-4.40
(2H,m), 4.75 (1H,d), 5.05 (1H,q), 5.70 (1H,d), 6.13

(2H,s), 6.89-6.99 (3H,m), 7.15-7.29 (3H,m), 7.36 (1H,d).
MS: 675 (M⁺). C₃₀H₃₇F₃N₂O₁₀S.

Example 89

5 Step1:

N-{(2*R*,3*S*)-3-Amino-4-[4-(benzyloxy)phenyl]-2-hydroxybutyl}-*N*-(5-cyano-2,2-dimethylpentyl)-1,3-benzodioxole-5-sulfonamide



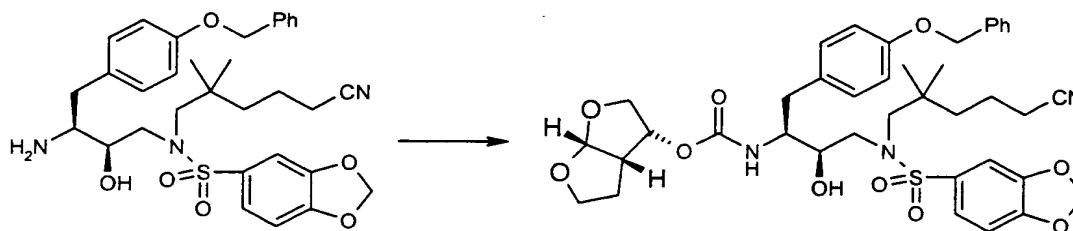
10

Treatment of tert-butyl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl) (5-cyano-2,2-dimethylpentyl)amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate with
15 trifluoroacetic acid/dichloromethane as previously described provided the title compound as a solid foam. ¹H
NMR (DMSO-*d*₆): δ 0.95 (6H, s), 1.15 (1H, br d), 1.2-1.6
(5H, m), 2.2 (1H, t), 2.42 (2H, br s), 2.6 (2H, br d),
2.9 (1H, d), 3.05 (1H, dd), 3.3 (1H, br s), 3.45 (2H, br
20 d), 4.62 (1H, s), 5.0 (2H, s), 6.15 (2H, s), 6.89 (2H,
d), 7.0 (1H, d), 7.06 (2H, d), 7.22-7.42 (7H, m); MS: 594
(MH⁺)

Step 2:

25 (3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl) (5-cyano-2,2-

dimethylpentyl) amino] -1- [4- (benzyloxy)benzyl] -2-
hydroxypropylcarbamate (305)

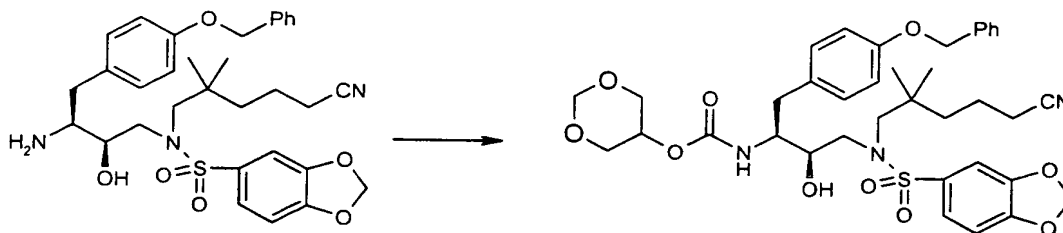


5 N- { (2R,3S) -3-Amino-4- [4- (benzyloxy) phenyl] -2-
hydroxybutyl} -N- (5-cyano-2,2-dimethylpentyl) -1,3-
benzodioxole-5-sulfonamide was treated with
[(3R,3aS,6aR) hexahydrofuro [2,3-*b*] furan-3-yl] [4-
nitrophenyl] carbonate, diisopropylethylamine and
10 acetonitrile as previously described to provide the title
compound as a solid foam. ¹H NMR (DMSO-*d*₆): δ 0.87 (3H,
s), 0.92 (3H, s), 1.10-1.13 (1H, m), 1.32-1.38 (3H, m),
1.5-1.6 (2H, m), 2.33 (1H, t), 2.4 (2H, t), 2.65-2.75
(2H, m), 2.7-2.9 (2H, m), 3.3-3.4 (3H, m), 3.5-3.6 (2H,
15 m), 3.65 (1H, t), 3.7 (1H, dd), 3.8 (1H, dd), 4.8 (1H,
dd), 5.0 (2H, s), 5.03 (1H, d), 5.45 (1H, d), 6.15 (2H,
s), 6.82 (2H, d), 7.03 (1H d), 7.04 (2H, d), 7.16 (1H,
d), 7.22 (1H, s), 7.26-7.40 (6H, m); MS: 750 (MH⁺);

20

Example 90

1,3-Dioxan-5-yl (1*S*,2*R*) -3- [(1,3-benzodioxol-5-
ylsulfonyl) (5-cyano-2,2-dimethylpentyl) amino] -1- [4-
(benzyloxy)benzyl] -2-hydroxypropylcarbamate (306)

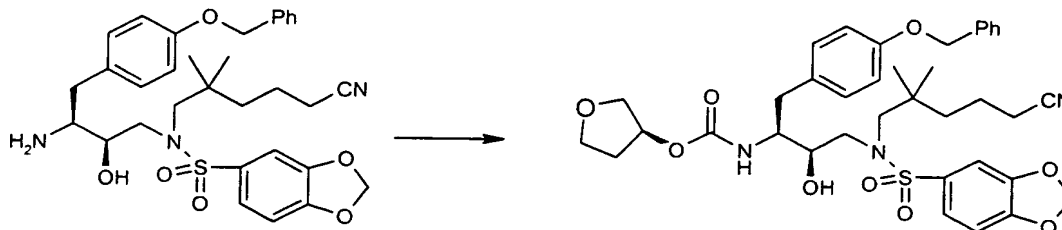


5 *N*-{(2*R*,3*S*)-3-Amino-4-[4-(benzyloxy)phenyl]-2-hydroxybutyl}-*N*-(5-cyano-2,2-dimethylpentyl)-1,3-benzodioxole-5-sulfonamide was treated with 1,3-dioxan-5-yl 4-nitrophenyl carbamate/diisopropylamine/acetonitrile as previously described to afford the title compound as a solid foam. ¹H NMR (DMSO-*d*₆): δ 0.88 (3H, s), 0.89 (3H, s), 1.3-1.4 (2H, m), 1.5-1.6 (2H, m), 2.4-2.5 (3H, m), 2.70-2.82 (2H, m), 2.95 (1H, dd), 3.3-3.4 (3H, m), 3.5 (1H, d), 3.65-3.75 (2H, m), 3.79 (1H, d), 3.89 (1H, d), 4.25 (1H, s), 4.65 (1H, d), 4.8 (1H, d), 4.97 (1H, d), 5.0 (2H, s), 6.15 (2H, s), 6.84 (2H, d), 7.0 (1H, d), 7.15 (2H, d), 7.2 (1H, d), 7.25 (1H, s), 7.26-7.40 (6H, m); MS: 724 (MH⁺)

15

Example 91

20 (3*S*)-Tetrahydro-3-furanyl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl) (5-cyano-2,2-dimethylpentyl) amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate (307)

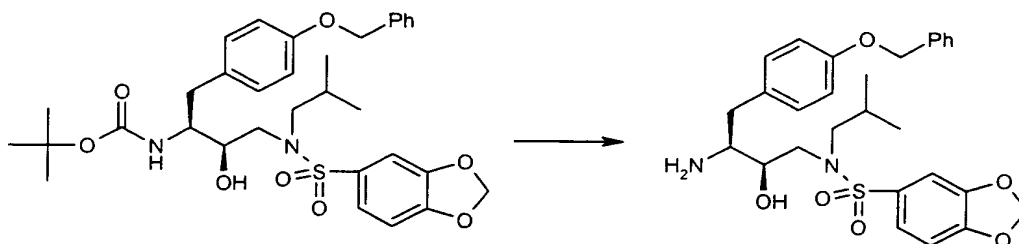


N-{(2*R*,3*S*)-3-Amino-4-[4-(benzyloxy)phenyl]-2-hydroxybutyl}-*N*-(5-cyano-2,2-dimethylpentyl)-1,3-benzodioxole-5-sulfonamide was treated with 1-({[(3*S*)-tetrahydro-3-furanyloxy]carbonyl}oxy)-2,5-pyrrolidinedione/diisopropylamine/acetonitrile as previously described to provide the title product as a solid foam. ¹H NMR (DMSO-*d*₆): δ 0.88 (3H, s), 0.90 (3H, s), 1.3 (2H, dd), 1.5-1.6 (2H, m), 1.7-1.8 (1H, m), 2.0-2.2 (1H, m), 2.37 (1H, t), 2.4 (2H, t), 2.75-2.85 (3H, m), 2.9 (1H, dd), 3.30-3.45 (3H, m), 3.55 (1H, dd), 3.65 (1H, dd), 3.7 (2H, dd), 4.9 (1H, s), 4.98 (1H, d), 5.0 (2H, s), 6.15 (2H, s), 6.8 (2H, d), 7.0 (2H, d), 7.06 (2H, d), 7.24 (1H, s), 7.25-7.42 (6H, m); MS: 708 (MH⁺);

Example 92

Step 1:

N-{(2*R*,3*S*)-3-Amino-4-[4-(benzyloxy)phenyl]-2-hydroxybutyl}-*N*-isobutyl-1,3-benzodioxole-5-sulfonamide (308)



Treatment of *tert*-butyl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-yl)(isobutyl)amino]-1-[4-(benzyloxy)benzyl]-2-

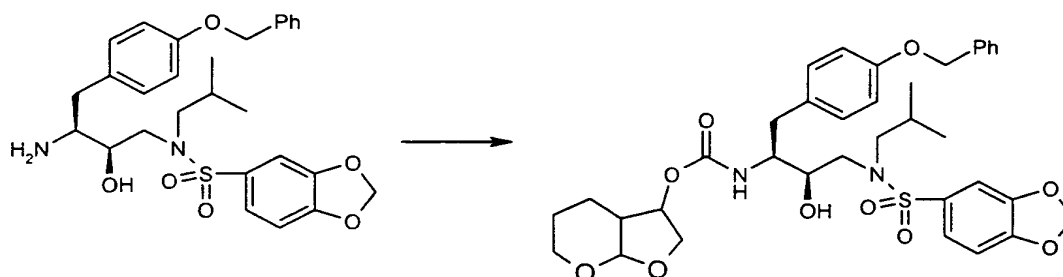
hydroxypropylcarbamate with trifluoroacetic acid as previously described afforded the title compound as a solid foam. ¹H NMR (DMSO-*d*₆): δ 0.90 (3H, d), 0.94 (3H,

d), 1.8-2.0 (2H, m), 2.3 (1H, dd), 2.70-2.85 (3H, m), 2.9-3.0 (2H, m), 3.2-3.3 (2H, m), 3.4-3.5 (2H, m), 4.7 (1H, d), 5.03 (2H, s), 6.18 (2H, s), 6.9 (2H, d), 7.02 (1H, d), 7.1 (2H, d), 7.24 (1H, s), 7.25-7.43 (5H, m);

5 MS: 527 (MH⁺)

Step 2:

Hexahydro-4*H*-furo[2,3-*b*]pyran-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate

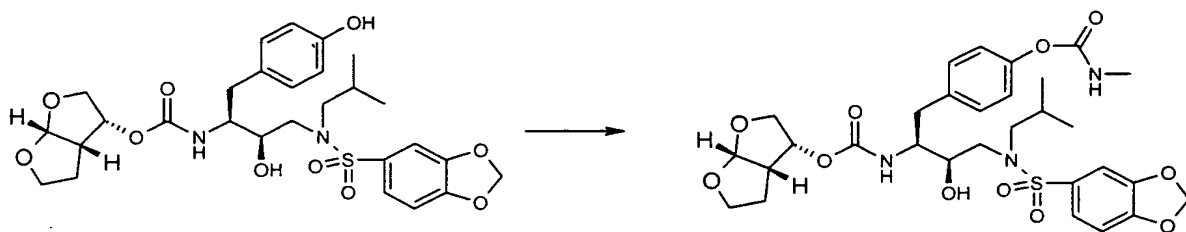


N-{(2*R*,3*S*)-3-Amino-4-[4-(benzyloxy)phenyl]-2-hydroxybutyl}-*N*-isobutyl-1,3-benzodioxole-5-sulfonamide was treated with hexahydro-4*H*-furo[2,3-*b*]pyran-3-yl 4-nitrophenyl carbonate/diisopropylethylamine/acetonitrile as previously described to afford after silica gel chromatography (dichloromethane/methanol, 49:1) the title diastereomers as a solid foams. Diastereomer A: ¹H NMR (DMSO-*d*₆): δ 0.76 (3H, d), 0.80 (3H, d), 1.2 (1H, br s), 1.6 (2H, br s), 1.9 (1H, br s), 2.1 (1H, br s), 2.4 (1H, br s), 2.75 (1H, dd), 2.8-3.0 (3H, m), 3.2-3.3 (3H, m), 3.4-3.6 (3H, m), 3.7 (1H, d), 4.0 (1H, t), 4.8-4.9 (2H, m), 5.0 (3H, br s), 6.15 (2H, s), 6.8 (2H, d), 7.0-7.2 (4H, m), 7.21-7.42 (7H, m); MS: 719 (M+23) Diastereomer B: ¹H NMR (DMSO-*d*₆): δ 0.76 (3H, d), 0.82 (3H, d), 1.0 (1H, br s), 1.3 (1H, br s), 1.5 (1H, br s), 1.95 (2H, br

s), 2.4 (1H, t), 2.7 (2H, td), 2.9 (1H, d), 2.97 (1H, dd), 3.2-3.3 (3H, m), 3.5 (1H, br s), 3.55 (1H, br s), 3.64 (2H, br s), 4.0 (1H, dd), 4.9 (1H, s), 4.95-5.05 (4H, m), 6.15 (2H, s), 6.8 (2H, d), 7.0-7.2 (4H, m), 7.22 (1H, s), 7.25-7.40 (6H, m); MS: 719 (M+23)

Example 93

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 10 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-(4-{[(methylamino)carbonyl]oxy}
 benzyl)propylcarbamate (309)

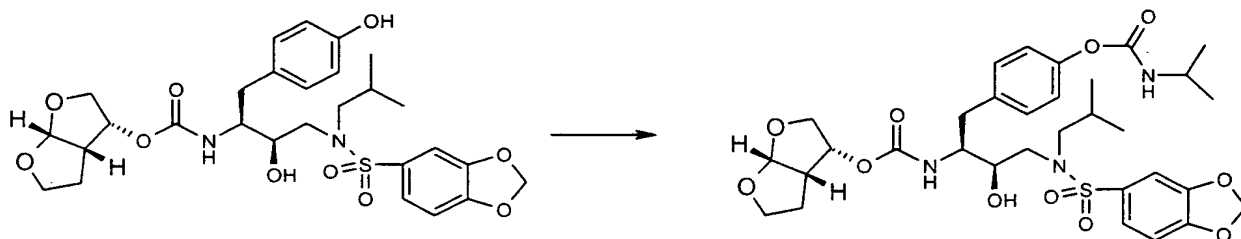


15 A mixture of (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl
 (1*S*,2*R*)-3-[(1,3-benzodioxol-5-
 ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-
 hydroxybenzyl)propylcarbamate (80 mg), methyl isocyanate
 20 (0.5 mL), dichloromethane (3 mL) and diisopropylamine
 (0.05 mL) was stirred at ambient temperature for 1 h.
 Solvent was evaporated and the residue was purified by
 chromatography (silica gel, hexanes/ethyl acetate, 3:1)
 to provide the title compound as a solid foam (57 mg). ¹H
 25 NMR (DMSO-*d*₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.17 (1H,
 dd), 1.30-1.45 (1H, m), 1.9-2.0 (1H, m), 2.42 (1H, t),
 2.6 (3H, d), 2.62-2.80 (3H, m), 2.99 (2H, dd), 3.15-3.20

(1H, m), 3.50-3.63 (4H, m), 3.7 (1H, t), 3.8 (1H, dd), 4.8 (1H, dd), 5.04 (1H, d), 5.5 (1H, d), 6.18 (2H, s), 6.95 (2H, d), 7.05 (1H, d), 7.15 (2H, d), 7.2 (1H, s), 7.25 (1H, d), 7.5 (1H, quartet), 8.08 (1H, d); MS: 650 (MH⁺).

Example 94

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 10 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-(4-{[(isopropylamino)carbonyl]oxy}
 benzyl)propylcarbamate (310)

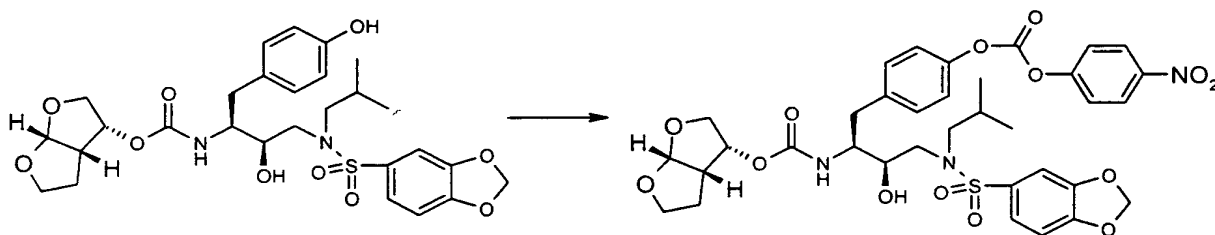


15 (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-(4-hydroxybenzyl)propylcarbamate was treated
 with isopropyl isocyanate/triethylamine/dichloromethane
 as described in Example 93 to provide the title compound
 20 as a solid foam. ¹H NMR (DMSO-d₆): δ 0.76 (3H, d), 0.82
 (3H, d), 0.98 (3H, d), 1.1 (3H, d), 1.35-1.42 (1H, m),
 1.90-1.95 (1H, m), 2.4 (1H, t), 2.65-2.80 (3H, m), 2.9-
 3.0 (2H, m), 3.3-3.4 (1H, m), 3.50-3.63 (6H, m), 3.7 (1H,
 t), 3.8 (1H, dd), 4.8 (1H, dt), 5.05 (1H, br d), 5.45
 25 (1H, d), 6.18 (2H, s), 6.92 (2H, d), 7.0 (1H, d), 7.18
 (2H, d), 7.22 (1H, s), 7.3 (2H, d), 7.58 (1H, d); MS: 678
 (MH⁺); C₃₂H₄₃N₃O₁₁S.

Example 95

Step 1:

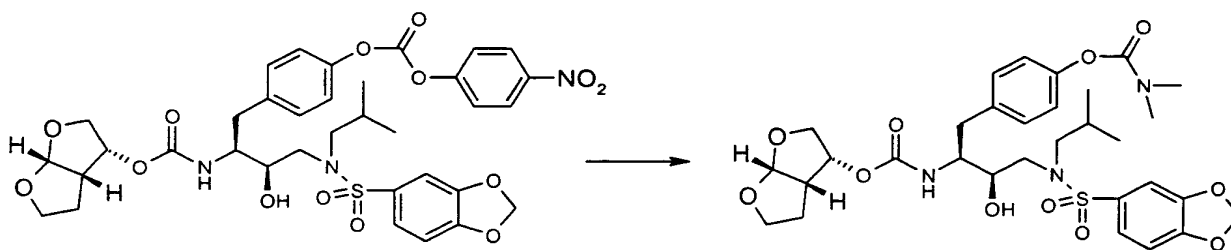
4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenyl 4-nitrophenyl carbonate



To a solution of (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-hydroxybenzyl)propylcarbamate (0.5 g, 0.84 mmol) in dichloromethane (4 mL) at 0°C was added pyridine (0.16 mL, 0.158 g, 2.0 mmol) and 4-nitrophenyl chloroformate (0.22 g, 1.09 mmol) and the mixture was stirred at ambient temperature for 1 h. Dichloromethane was added and the mixture was washed with 15% citric acid/water (2X), saturated sodium bicarbonate/water, dried (sodium sulfate), evaporated, and purified by chromatography (silica gel, dichloromethane/methanol, 49:1) to provide the title compound as a solid foam (0.5 g, 79% yield). ¹H NMR (DMSO-*d*₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.16 (1H, dd), 1.3-1.4 (1H, m), 1.9-2.0 (1H, m), 2.42 (1H, t), 2.65-2.80 (3H, m), 3.0 (2H, dd), 3.3-3.4 (1H, m), 3.5-3.6 (4H, m), 3.7 (1H, t), 3.8 (1H, dd), 4.8 (1H, dt), 5.1 (1H, d), 5.5 (1H, d), 6.18 (2H, s), 7.05 (1H, d), 7.2-7.4 (7H, m), 7.7 (2H, d), 8.35 (2H, d)

Step 2:

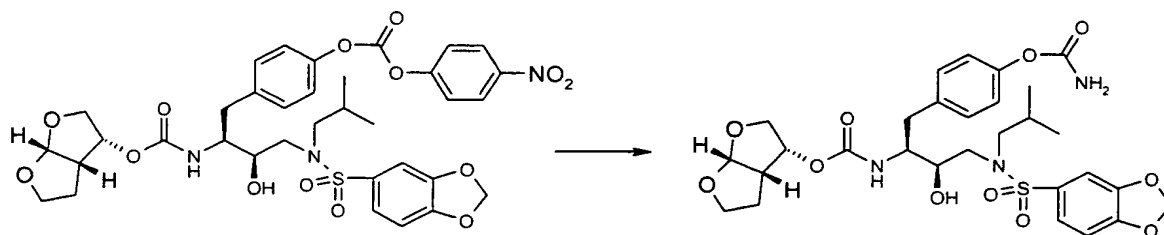
(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-(4-
 5 {[(dimethylamino)carbonyl]oxy}benzyl)-2-
 hydroxypropylcarbamate (311)



To a solution of 4-((2S,3R)-2-({[(3R,3aS,6aR)-
 10 Hexahydrofuro[2,3-b]furan-3-yloxy]carbonyl}amino)-4-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-3-
 hydroxybutyl}phenyl 4-nitrophenyl carbonate (60 mg) in
 tetrahydrofuran (0.5 mL) was added 2M
 dimethylamine/tetrahydrofuran (0.5 mL) and the mixture
 15 was stirred at ambient temperature for 30 min. Ethyl
 acetate was added and the mixture was washed with
 saturated sodium bicarbonate/water (4X), dried (sodium
 sulfate), evaporated, and purified by chromatography
 (silica gel, hexanes/ethyl acetate, 3:7) to provide the
 20 title compound as a solid foam. ¹H NMR (DMSO-d₆): δ 0.76
 (3H, d), 0.82 (3H, d), 1.1-1.2 (1H, m), 1.2 (1H, br
 quintuplet), 1.9-2.0 (1H, m), 2.2 (1H, t), 2.6-2.8 (3H,
 m), 2.81 (3H, s), 2.9 (3H, s), 3.0 (1H, br s), 3.2-3.3
 (2H, m), 3.4-3.6 (4H, m), 3.7 (1H, t), 3.8 (1H, dd), 4.8
 25 (1H, dt), 5.0 (1H, br s), 5.5 (1H, d), 6.1 (2H, s), 6.9
 (2H, d), 7.0 (1H, d), 7.15 (2H, d), 7.19 (1H, s), 7.22
 (2H, br d); MS: 664 (MH⁺); C₃₁H₄₁N₃O₁₁S.

Example 96A

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-{4-
[(aminocarbonyl)oxy]benzyl}-3-[(1,3-benzodioxol-5-
ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate
5 (312)

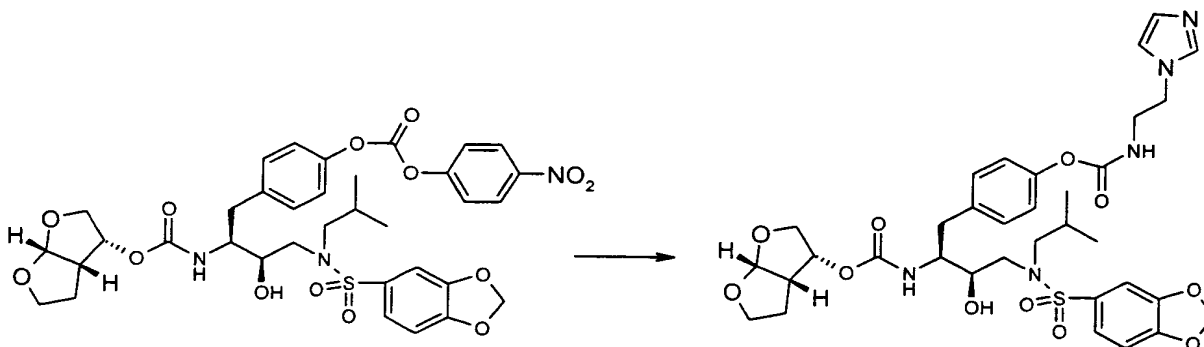


4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-
yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-
10 ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenyl 4-
nitrophenyl carbonate was treated with concentrated
ammonium hydroxide as described in Example 311 to afford
the title compound as a white solid. ¹H NMR (DMSO-*d*₆): δ
0.78 (3H, d), 0.86 (3H, d), 1.2 (1H, dd), 1.4 (1H, br
15 quintuplet), 1.85-1.95 (1H, m), 2.2 (1H, t), 2.65-2.80
(3H, m), 2.9-3.0 (2H, m), 3.25-3.30 (1H, m), 3.5-3.6 (4H,
m), 3.7 (1H, t), 3.8 (1H, dd), 4.8 (1H, dt), 5.0 (1H, d),
5.4 (1H, d), 6.18 (2H, d), 6.8 (2H, br d), 6.9 (2H, d),
7.0 (1H, d), 7.15 (2H, d), 7.2 (1H, s), 7.25-7.30 (2H,
20 m); MS: 636 (MH⁺).

Example 96B

25 (3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-

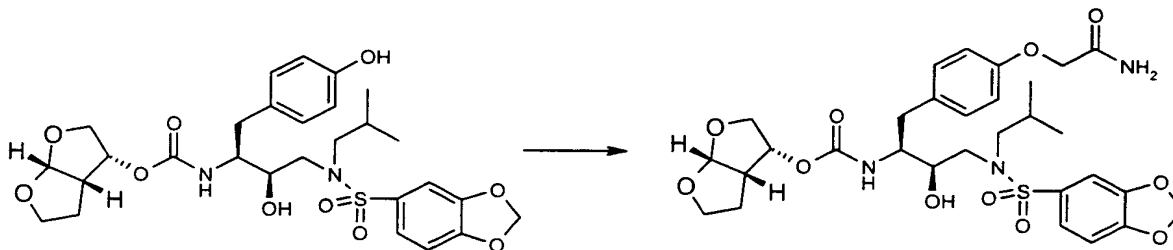
hydroxy-1-{4-[(2-(1H-imidazol-1-yl)ethyl)amino]carbonyloxy]benzyl}propylcarbamate (313)



- 5 To a solution of 4-{(2S,3R)-2-([(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yloxy]carbonyl)amino)-4-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenyl 4-nitrophenyl carbonate (50 mg, 0.07 mmol) in 1,4-dioxane (1 mL) was added 2-(1H-imidazol-1-yl)ethanamine (50 mg, 0.45 mmol, *Synthetic Communications* 1991, 21, 535-544) and the mixture was stirred at ambient temperature for 30 min. The mixture was diluted with ethyl acetate and washed with saturated sodium bicarbonate/water (4X), dried (sodium sulfate),
- 15 evaporated and purified by chromatography (silica gel, dichloromethane/2M ethanolic ammonia, 93:7) to provide the title compound as a white solid (25 mg). ¹H NMR (DMSO-d₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.2 (1H, dd), 1.4 (1H, quintuplet), 1.9-2.0 (1H, m), 2.4 (1H, t), 2.55-2.80 (3H, m), 2.9-3.0 (2H, m), 3.3-3.4 (3H, m), 3.5-3.6 (4H, m), 3.7 (1H, t), 3.8 (1H, dd), 4.03-4.07 (2H, m), 4.8 (1H, dt), 5.05 (1H, d), 5.5 (1H, d), 6.18 (2H, s), 6.87 (1H, s), 6.9 (2H, d), 7.05 (1H, d), 7.15 (1H, s), 7.2 (2H, d), 7.23 (1H, s), 7.25-7.30 (2H, m), 7.6 (1H, s), 7.8 (1H, t); MS: 730 (MH⁺)
- 25

Example 97

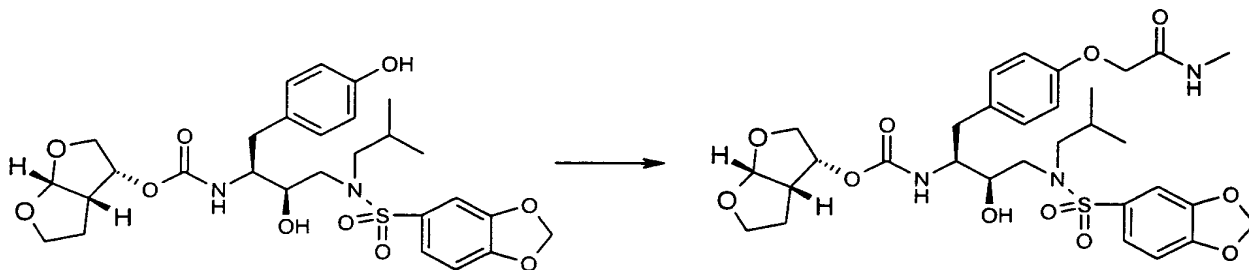
(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-1-[4-(2-amino-2-oxoethoxy)benzyl]-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate (314)



To a solution of (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-hydroxybenzyl)propylcarbamate (47 mg, 0.08 mmol) in anhydrous dimethylformamide (2 mL) was added cesium carbonate (78 mg, 0.24 mmol) and 2-bromoacetamide (22 mg, 0.16 mmol). After 1 h at ambient temperature, the mixture was diluted with ethyl acetate, washed with water (3X), brine, dried (sodium sulfate), and evaporated. The residue was purified by chromatography (silica gel, dichloromethane/methanol, 97:3) to afford the title product as a white solid (45 mg). ¹H NMR (DMSO-d₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.2 (1H, dd), 1.4 (1H, quintuplet), 1.9-2.0 (1H, m), 2.35 (1H, t), 2.6-2.8 (3H, m), 2.9 (1H, d), 2.95 (1H, dd), 3.25-3.30 (1H, m), 3.45 (1H, br s), 3.5-3.6 (3H, m), 3.7 (1H, t), 3.8 (1H, dd), 4.3 (2H, s), 4.8 (1H, dt), 5.0 (1H, d), 5.5 (1H, d), 6.18 (2H, s), 6.8 (2H, d), 7.0-7.15 (3H, m), 7.22 (1H, s), 7.24 (1H, d), 7.26 (1H, d), 7.32 (1H, s), 7.42 (1H, s); MS: 650 (MH⁺)

Example 98

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-{4-[2-(methylamino)-2-oxoethoxy]benzyl}
 5 propylcarbamate (315)

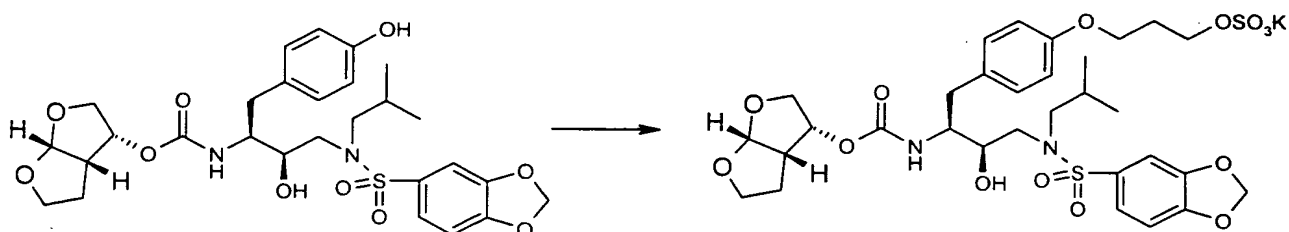


To a solution of (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-
 3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-
 10 ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-
 hydroxybenzyl)propylcarbamate (60 mg, 0.1 mmol) in
 anhydrous tetrahydrofuran (1.5 mL) was added 60% sodium
 hydride/mineral oil dispersion (4 mg, 0.1 mmol). The
 mixture was stirred for 10 min at ambient temperature
 15 under nitrogen atmosphere and a solution of *N*-methyl-2-
 bromoacetamide (17 mg, 0.11 mmol) in anhydrous
 tetrahydrofuran (0.5 mL) was added. After 2 h, an
 additional portion of 60% sodium hydride/mineral oil
 dispersion (4 mg, 0.1 mmol) was added. After 15 min,
 20 acetic acid (0.1 mL) was added and the mixture was
 diluted with ethyl acetate, washed with saturated sodium
 bicarbonate/water, dried (sodium sulfate), evaporated,
 and purified by chromatography (silica gel, ethyl
 acetate) to provide the title compound as a white solid
 25 (10 mg). ¹H NMR (DMSO-*d*₆): δ 0.78 (3H, d), 0.82 (3H, d),
 1.15-1.22 9 (1H, m), 1.2 (1H, quintuplet), 1.9-2.0 (1H,
 m), 2.35 (1H, t), 2.6 (3H, d), 2.65-2.80 (3H, m), 2.9

(1H, d), 2.95 (1H, dd), 3.2-3.3 (1H, m), 3.4-3.5 (1H, m),
 3.5-3.6 (3H, m), 3.7 (1H, t), 3.8 (1H, dd), 4.4 (2H, s),
 4.8 (1H, dt), 5.0 (1H, d), 5.5 (1H, d), 6.2 (2H, s), 6.8
 (2H, d), 7.0-7.2 (3H, m), 7.22 (1H, s), 7.25 (1H, d), 7.3
 5 (1H, d), 8.0 (1H, br s); MS: 664 (MH⁺);

Example 99

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 10 hydroxy-1-{4-[3-(sulfoxy)propoxy]benzyl}propylcarbamate
 Potassium Salt (316)



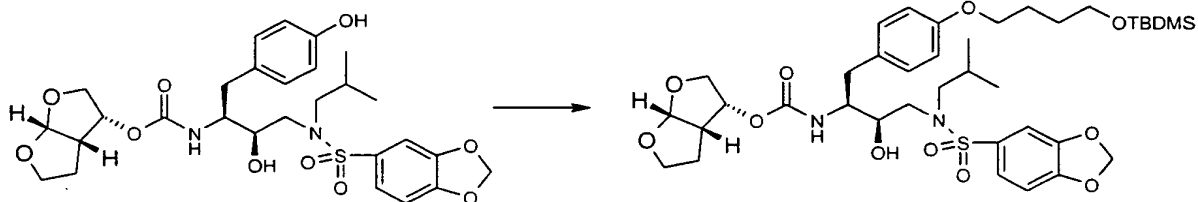
A mixture of (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl
 (1S,2R)-3-[(1,3-benzodioxol-5-
 15 ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-
 hydroxybenzyl)propylcarbamate (35 mg, 0.06 mmol),
 potassium carbonate (8.1 mg, 0.06 mmol), 1,3,2-
 dioxathiane 2,2-dioxide (8.9 mg, 0.065 mmol, *J. Am. Chem.*
Soc. **1988**, *110*, 7538-7539) and acetonitrile was heated
 20 at 82°C under nitrogen atmosphere for 36 h and filtered
 while hot. The filtrate was allowed to cool to ambient
 temperature for 18 h and the resulting solid was filtered
 and washed with diethyl ether to provide the title
 compound as a white solid (25 mg). ¹H NMR (DMSO-d₆): δ
 25 0.78 (3H, d), 0.82 (3H, d), 1.25 (1H, dd), 1.4 (1H,
 quintuplet), 1.9-2.0 (3H, m), 2.2 (1H, t), 2.7-2.8 (3H,
 m), 2.9 (1H, d), 3.0 (1H, dd), 3.25-3.00 (1H, m), 3.4-3.5

(1H, m), 3.50-3.65 (3H, m), 3.7 (1H, t), 3.75-3.82 (3H, m), 3.9 (1H, t), 4.08 (1H, dd), 4.8 (1H, dt), 4.97 (1H, d), 5.5 (1H, d), 6.18 (2H, s), 6.78 (2H, d), 7.0-7.1 (3H, m), 7.19 (1H, d), 7.20 (1H, s), 7.30 (1H, d); MS: 730 (MH⁺)

Example 100

Step 1:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 10 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(4-
 {[tert-butyl(dimethyl)silyl]oxy}butoxy)benzyl]-2-
 hydroxypropylcarbamate

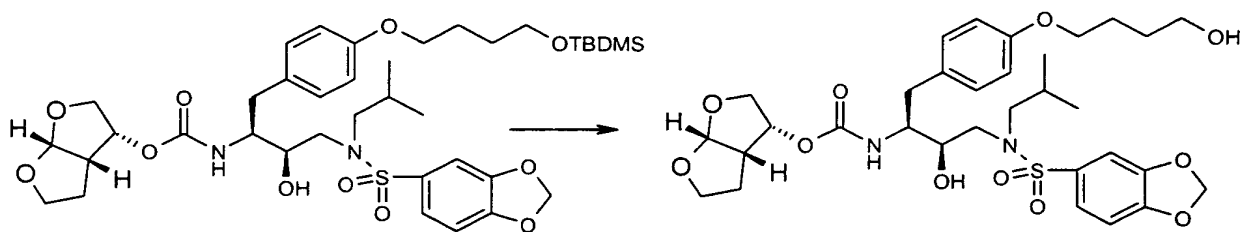


15 To a solution of (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-
 3-yl (1S,2R)-3-[(1,3-benzodioxol-5-
 ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-
 hydroxybenzyl)propylcarbamate (0.6 g, 1.01 mmol),
 triphenyl phosphine (0.4 g, 1.52 mmol), and 4-{[tert-
 20 butyl(dimethyl)silyl]oxy}-1-butanol (0.31 g, 1.52 mmol,
J. Org. Chem. **1986**, *51*, 3388-3390) in anhydrous
 dichloromethane (9 mL) at 0°C under nitrogen atmosphere
 was slowly added a solution of diisopropyl
 azodicarboxylate (0.29 mL, 0.3 g, 1.52 mmol) in anhydrous
 25 dichloromethane (2 mL) and the mixture was stirred at
 ambient temperature for 2 h. Solvent was evaporated and
 the residue was purified by chromatography (silica gel,

hexanes/ethyl acetate, 1:1) to provide the title compound as a solid foam (0.39 g). ¹H NMR (DMSO-d₆): δ 0.0 (6H, s), 0.78 (3H, d), 0.80 (3H, d), 0.82 (9H, s), 1.22 (1H, dd), 1.4 (1H, quintuplet), 1.55 (2H, quintuplet), 1.7 (2H, quintuplet), 1.8-1.9 (1H, m), 2.35 (1H, t), 2.65-2.80 (3H, m), 2.9 (1H, d), 3.0 (1H, dd), 3.2-3.3 (2H, m), 3.4-3.5 (1H, m), 3.52-3.62 (4H, m), 3.7 (1H, t), 3.8 (1H, dd), 3.9 (2H, t), 4.8 (1H, dt), 5.0 (1H, br s), 5.5 (1H, d), 6.18 (2H, s), 6.77 (2H, d), 7.0-7.1 (3H, m), 7.19 (1H, d), 7.21 (1H, s), 7.3 (1H, d); MS: 801 (M+23); C₃₈H₅₈N₂O₁₁SSi.

Step2:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-[4-(4-hydroxybutoxy)benzyl]propylcarbamate (317)



20

To a solution of (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(4-{[tert-butyl(dimethyl)silyl]oxy}butoxy)benzyl]-2-hydroxypropylcarbamate (0.38 g, 0.48 mmol) in tetrahydrofuran (5 mL) at 0°C was added a 1:1 mixture of 1M tetra-*n*-butyl ammonium fluoride/tetrahydrofuran and acetic acid (1.2 mL) and the mixture was stirred at

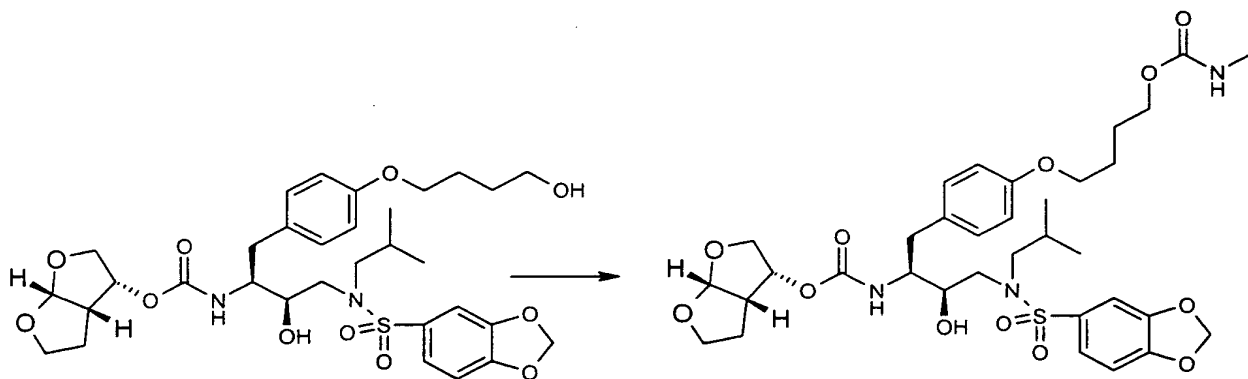
25

ambient temperature for 18 h. The mixture was diluted with ethyl acetate and washed with water (4X), dried (sodium sulfate) and evaporated. The resulting solid was triturated with diethyl ether and filtered to afford the
 5 title compound as a white solid (0.24 g). ¹H NMR (DMSO-
_d₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.2 (1H, dd), 1.38 (1H, quintuplet), 1.5 (2H, quintuplet), 1.65 (2H, quintuplet), 1.9-2.0 (1H, m), 2.18 (1H, t), 2.7-2.8 (3H, m), 2.9 (1H, d), 2.97 (1H, dd), 3.20-3.25 (1H, m), 3.38 (2H, t), 3.4-
 10 3.5 (1H, m), 3.52-3.62 (3H, m), 3.7 (1H, t), 3.8 (1H, dd), 3.93 (2H, t), 4.2 (1H, br s), 4.8 (1H, dt), 5.0 (1H, d), 5.5 (1H, d), 6.18 (2H, s), 6.78 (2H, d), 7.0-7.1 (3H, m), 7.2 (1H, d), 7.26 (1H, s), 7.3 (1H, d); MS: 665 (MH⁺)

15

Example 101

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-(4-{[(methylamino)carbonyl]oxy}butoxy)
 20 benzyl]propylcarbamate (318)



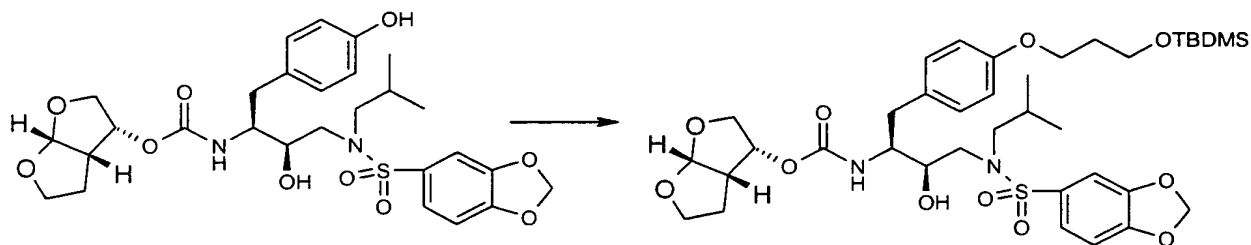
(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 25 hydroxy-1-[4-(4-hydroxybutoxy)benzyl]propylcarbamate was

treated with methyl isocyanate/dichloromethane as described earlier to afford the title compound as a solid foam. ¹H NMR (DMSO-d₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.2 (1H, dd), 1.4 (1H, quintuplet), 1.6-1.8 (4H, m), 1.9-2.0 (1H, m), 2.38 (1H, t), 2.5 (3H, d), 2.7-2.8 (3H, m), 2.9 (1H, d), 2.98 (1H, dd), 3.3-3.4 (1H, m), 3.4-3.5 (1H, m), 3.5-3.6 (3H, m), 3.7 (1H, t), 3.8 (1H, dd), 3.9 (2H, t), 3.95 (2H, t), 4.8 (1H, dt), 5.0 (1H, br s), 5.5 (1H, d), 6.18 (2H, s), 6.78 (2H, d), 6.9 (1H, br s), 7.0-7.1 (3H, m), 7.2 (1H, d), 7.25 (1H, s), 7.3 (1H, d); MS: 744 (M+23); C₃₄H₄₇N₃O₁₂S.

Example 102

Step 1:

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(3-{[*tert*-butyl(dimethyl)silyl]oxy}propoxy)benzyl]-2-hydroxypropylcarbamate

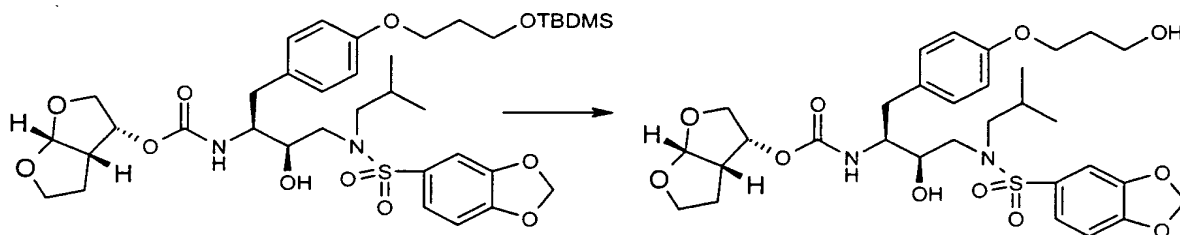


(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-hydroxybenzyl)propylcarbamate was treated with 3-{[*tert*-butyl(dimethyl)silyl]oxy}-1-propanol (*J. Org. Chem.* **1986**, *51*, 3388-3390)/triphenyl phosphine/diisopropyl azodicarboxylate as described in

step a (Example 213) to provide the title compound as solid foam. ¹H NMR (DMSO-*d*₆): δ 0.0 (6H, s), 0.78 (3H, d), 0.8 (9H, s), 0.82 (3H, d), 1.2 (1H, dd), 1.4 (1H, quintuplet), 1.8 (2H, quintuplet), 1.9-2.0 (1H, m), 2.38 (1H, dd), 2.6-2.8 (3H, m), 2.9 (1H, d), 2.98 (1H, dd), 3.2-3.3 (2H, m), 3.35-3.45 (1H, m), 3.5-3.6 (3H, m), 3.7 (2H, t), 3.8 (1H, dd), 3.9 (2H, t), 4.8 (1H, dt), 5.0 (1H, d), 5.5 (1H, d), 6.18 (2H, s), 6.75 (2H, d), 7.0-7.1 (3H, m), 7.18 (1H, d), 7.2 (1H, s), 7.28 (1H, d); MS: 765 (MH⁺)

Step 2:

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-[4-(3-hydroxypropoxy)benzyl]propylcarbamate (319)



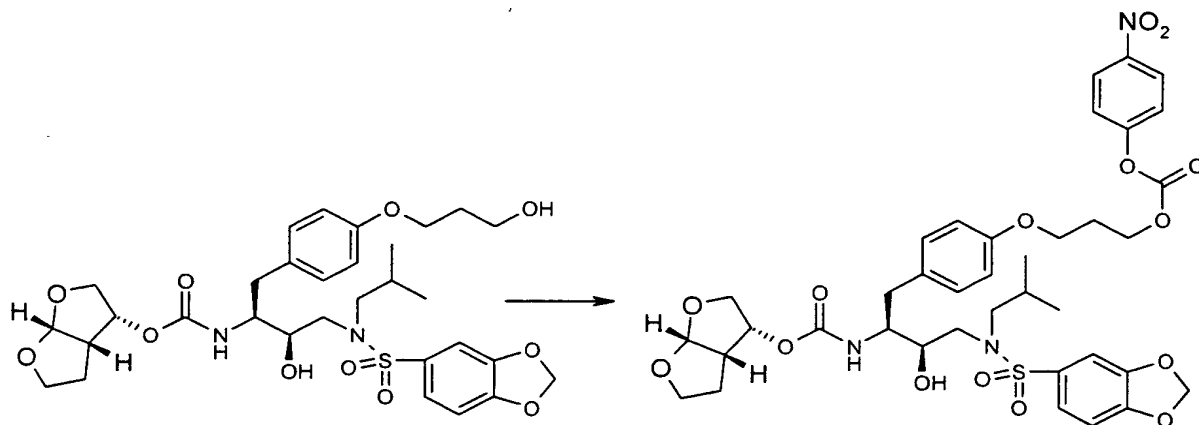
(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(3-{[*tert*-butyl(dimethyl)silyl]oxy}propoxy)benzyl]-2-hydroxypropylcarbamate was treated with 1M tetra-*n*-butyl ammonium fluoride/tetrahydrofuran/acetic acid as described in step b (Example 213) to afford the title compound as a white solid. ¹H NMR (DMSO-*d*₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.2 (1H, dd), 1.38 (1H, quintuplet), 1.8 (2H, quintuplet), 1.9-2.0 (1H, m), 2.38 (1H, t), 2.6-

2.8 (3H, m), 2.9 (1H, d), 2.96 (1H, dd), 3.3 (1H, br s),
3.4-3.6 (6H, m), 3.7 (1H, t), 3.8 (1H, dd), 3.9 (2H, t),
4.5 (1H, t), 4.8 (1H, dt), 5.0 (1H, d), 5.5 (1H, d), 6.18
(2H, s), 6.78 (2H, d), 7.0-7.1 (3H, m), 7.2 (1H, d), 7.25
5 (1H, s), 7.3 (1H, d); MS: 651 (MH⁺)

Example 103

Step 1:

10 3-(4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-
b]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-
ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenoxy)propyl
4-nitrophenyl carbonate



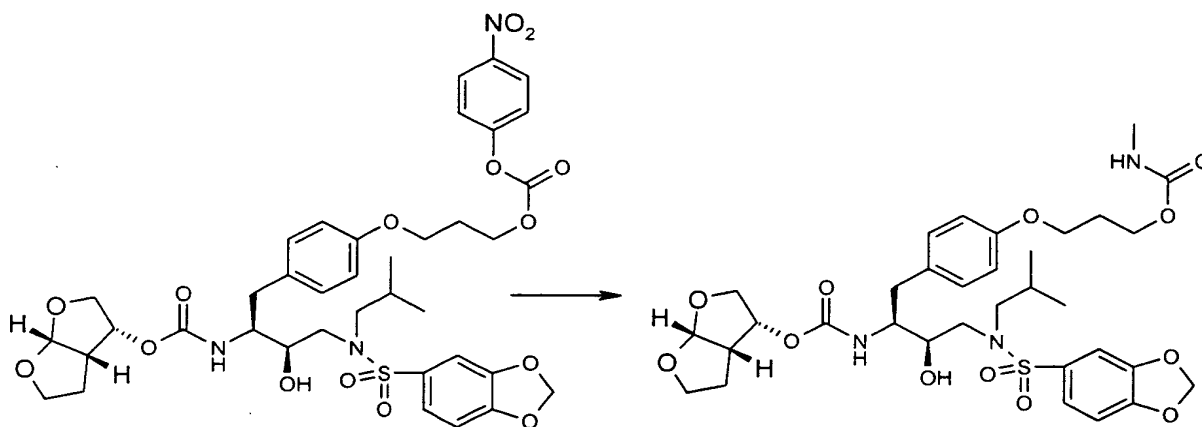
15

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-[4-(3-hydroxypropoxy)benzyl]propylcarbamate was
20 treated with 4-nitrophenyl chloroformate as described
previously to provide the title compound as a solid foam.
¹H NMR (DMSO-*d*₆): δ 0.86 (3H, d), 0.92 (3H, d), 1.2 (1H,
dd), 1.38 (1H, quintuplet), 1.9-2.0 (1H, m), 2.05-2.15

(2H, m), 2.2 (1H, t), 2.6-2.8 (3H, m), 2.9 (1H, d), 2.96 (1H, dd), 3.2-3.3 (2H, m), 3.4-3.5 (1H, m), 3.5-3.6 (2H, m), 3.7 (1H, t), 3.8 (1H, dd), 4.0 (2H, t), 4.37 (2H, t), 4.8 (1H, dt), 5.0 (1H, d), 5.45 (1H, d), 6.18 (2H, s),
5 6.78 (2H, d), 7.0-7.1 (3H, m), 7.20-7.25 (2H, m), 7.3 (1H, dd), 7.56 (2H, d), 8.25 (2H, d); MS: 816 (MH⁺)

Step 2:

10 (3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-[4-(3-{[(methylamino)carbonyl]oxy}propoxy)
benzyl]propylcarbamate (320)



15

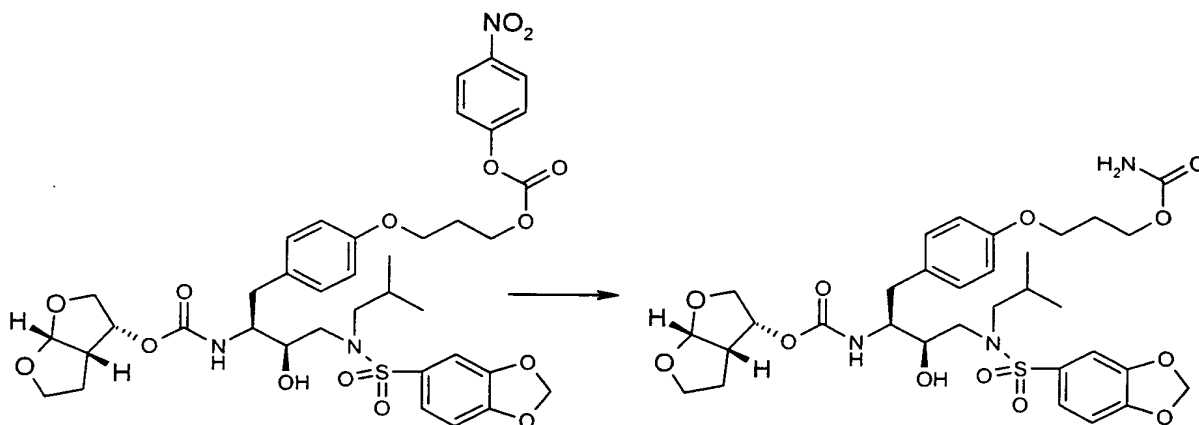
3-(4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-
b]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-
ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenoxy)propyl
4-nitrophenyl carbonate was treated with 2M

20 methylamine/tetrahydrofuran as described above to provide
the title compound as a white solid. ¹H NMR (DMSO-*d*₆): δ
0.78 (3H, d), 0.84 (3H, d), 1.1-1.2 (1H, m), 1.2 (1H, br
quintuplet), 1.9-2.0 (3H, m), 2.17 (1H, t), 2.45 (3H, d),

2.6-2.8 (3H, m), 2.83-3.00 (2H, m), 3.4-3.6 (5H, m), 3.7 (1H, br s), 3.8 (1H, br s), 3.9 (2H, br s), 4.0 (2H, br s), 4.8-4.9 (1H, m), 5.0 (1H, d), 5.5 (1H, d), 6.17 (2H, s), 6.7-6.8 (2H, m), 6.9 (1H, br s), 7.0-7.1 (3H, m),
5 7.2-7.4 (3H, m); MS: 708 (MH⁺);

Example 104

(3R,3aS,6aR) -Hexahydrofuro[2,3-b]furan-3-yl (1S,2R) -1-(4-
10 {3-[(aminocarbonyl)oxy]propoxy}benzyl)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxypropylcarbamate (321)



15 3-(4-{(2S,3R)-2-({[(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenoxy)propyl
4-nitrophenyl carbonate was treated with concentrated ammonium hydroxide as described above to afford the title
20 compound as a white solid. ¹H NMR (DMSO-d₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.2-1.3 (1H, m), 1.4 (1H, quintuplet), 1.9-2.0 (3H, m), 2.2 (1H, t), 2.6-2.8 (3H, m), 2.9 (1H, d), 2.97 (1H, dd), 3.2-3.3 (1H, m), 3.4-3.6 (4H, m), 3.7 (1H, t), 3.8 (1H, dd), 3.9 (2H, t), 4.0 (2H, t), 4.8 (1H,

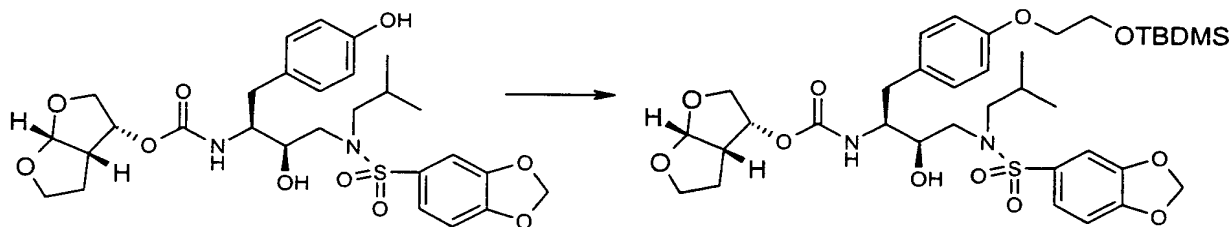
dt), 5.0 (1H, d), 5.5 (1H, d), 6.17 (2H, s), 6.4 (2H, br s), 6.78 (2H, d), 7.0-7.1 (3H, m), 7.2-7.3 (3H, m); MS: 694 (MH⁺)

5

Example 105

Step 1:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-(1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(2-
 10 { [tert-butyl(dimethyl)silyl]oxy}ethoxy)benzyl]-2-
 hydroxypropylcarbamate

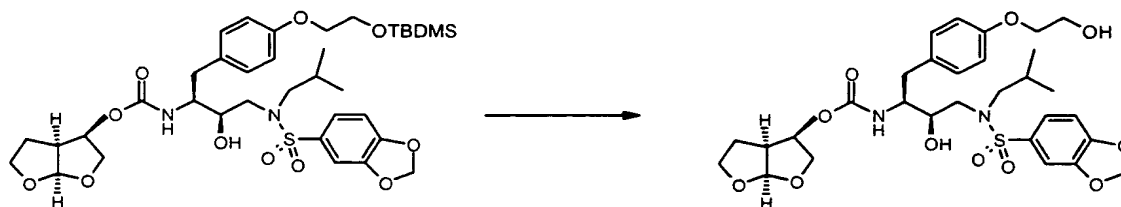


(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 15 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-(4-hydroxybenzyl)propylcarbamate was treated
 with 2-{ [tert-butyl(dimethyl)silyl]oxy}-1-ethanol (J.
 Org. Chem. 1986, 51, 3388-3390)/triphenyl
 phosphine/diisopropyl azodicarboxylate as described above
 20 to provide the title compound as solid foam. ¹H NMR (DMSO-
 d₆): 0.0 (6H, s), 0.78 (3H, d), 0.82 (3H, d), 0.84 (9H,
 s), 1.2 (1H, dd), 1.37 (1H, quintuplet), 1.9-2.0 (1H, m),
 2.37 (1H, dd), 2.6-2.8 (2H, m), 2.9 (1H, d), 2.98 (1H,
 dd), 3.2-3.3 (1H m), 3.4-3.5 (1H, m), 3.5-3.6 (4H, m),
 25 3.7 (1H, t), 3.8 (1H, dd), 3.88 (2H, dd), 3.93 (2H, dd),
 4.8 (1H, dt), 5.0 (1H, d), 5.45 (1H, d), 6.18 (2H, s),
 6.78 (2H, d), 7.0-7.1 (3H, m), 7.18 (1H, d), 7.2 (1H, s),
 7.28 (1H, d); MS: 751 (MH⁺)

000050 "494T6560

Step 2:

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 5 hydroxy-1-[4-(2-hydroxyethoxy)benzyl]propylcarbamate
 (322)



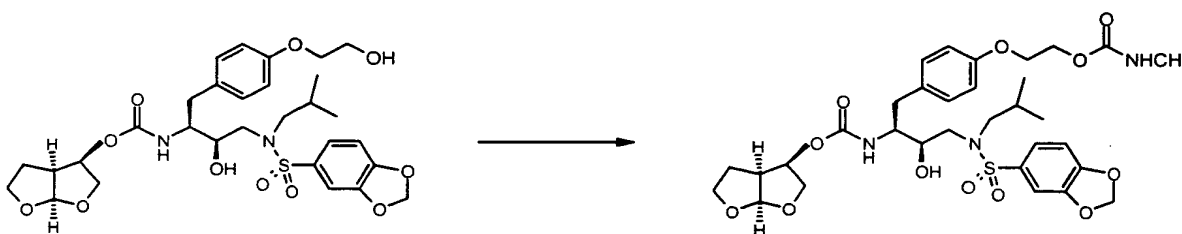
10

To a stirred solution of (3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(2-{[tert-butyl(dimethyl)silyl]oxy}ethoxy)benzyl]-2-
 15 hydroxypropylcarbamate (160mg, 0.21 mmol) in tetrahydrofuran (3 mL) at 5 °C was added a mixture of tetrabutylammonium fluoride (0.3 mL, 1M in tetrahydrofuran) and glacial acetic acid (0.3 mL) over 2 minutes. The reaction was allowed to warm to ambient
 20 temperature and stirred for 18 hours. The mixture was diluted with ethyl acetate (50 mL), washed with water (25 mL) followed by saturated sodium bicarbonate (20 mL) and then brine (20 mL), dried (magnesium sulfate) and concentrated *in vacuo* to afford the title compound (135
 25 mg, quant.) as a white solid. ¹H NMR (DMSO-*d*₆): δ 0.78 (6H, dd), 1.17-1.42 (2H, m), 1.92 (1H, m), 2.33 (1H, t), 2.64-2.78 (3H, m), 2.89-3.02 (2H, m), 3.24-3.29 (1H, m), 3.52-3.73 (7H, m), 3.78-3.82 (1H, m), 3.86 (2H, t), 4.78-

4.82 (2H, m), 4.99 (1H, d), 5.46 (1H, d), 6.12 (2H, s),
6.73 (2H, d), 7.02-7.28 (6H, m); MS: 637 (MH⁺)

Example 106

- 5 (3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-[4-(2-{[(methylamino)carbonyl]oxy}ethoxy)
benzyl]propylcarbamate (323)

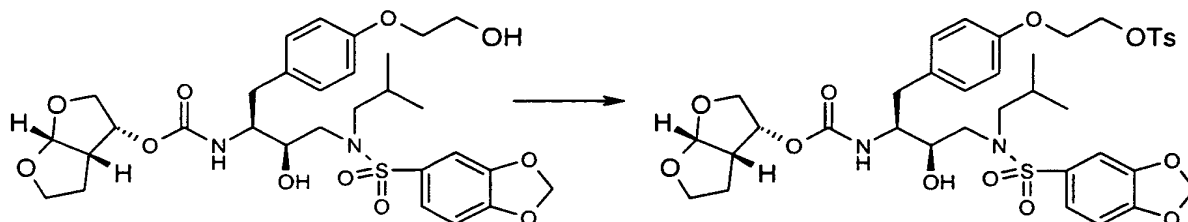


- 10 Treatment of (3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl
(1*S*,2*R*)-3-[(1,3-benzodioxol-5-
ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-[4-(2-
hydroxyethoxy)benzyl]propylcarbamate with methyl
isocyanate in dichloromethane as described earlier
15 afforded the title compound as a white foam in 52% yield.
¹H NMR (DMSO-*d*₆): δ 0.78 (6H, dd), 1.17-1.42 (2H, m), 1.92
(1H, m), 2.33 (1H, t), 2.60 (3H, d), 2.64-2.78 (3H, m),
2.88-2.98 (2H, m), 3.24-3.29 (1H, m), 3.40-3.60 (4H, m),
3.70 (1H, t), 3.78-3.82 (1H, m), 3.97-4.04 (2H, m), 4.19
20 (2H, t), 4.80 (1H, q), 4.99 (1H, d), 5.46 (1H, d), 6.12
(2H, d), 6.74 (2H, d), 7.02-7.28 (6H, m), 8.07 (1H, m);
MS: 694 (MH⁺)

Example 107

- 25 Step 1:

2-(4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-
b]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-
ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenoxy)ethyl
4-methylbenzenesulfonate

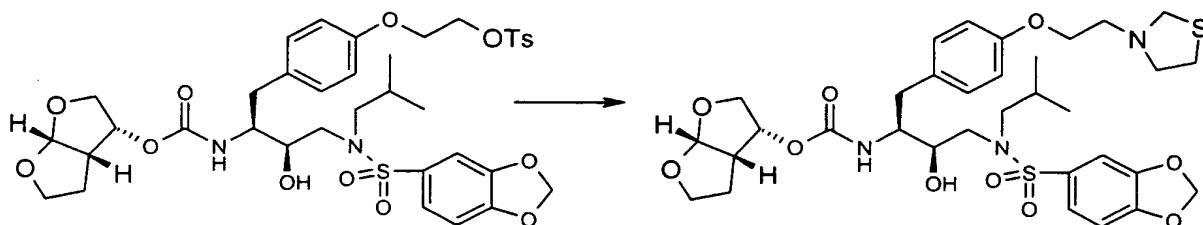


5

To a solution of (3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-[4-(2-
10 hydroxyethoxy)benzyl]propylcarbamate (0.11 g, 0.18 mmol), triethyl amine (0.055 mL, 40 mg, 0.39 mmol), and 4-dimethylaminopyridine (2 mg), in dichloromethane at 0°C was added *p*-toluenesulfonyl chloride (38 mg, 0.2 mmol) and the mixture was stirred at ambient temperature for 4
15 h. Solvent was evaporated and the residue was purified by chromatography (silica gel, dichloromethane/methanol, 98:2) to provide the title compound as a solid foam (0.105 g). ¹H NMR (DMSO-*d*₆): δ 0.78 (3H, d), 0.82 (3H, d), 1.2 (1H, dd), 1.37 (1H, quintuplet), 1.9-2.0 (1H, m), 2.3
20 (1H, dd), 2.38 (3H, s), 2.6-2.8 (2H, m), 2.9 (1H, d), 2.97 (1H, dd), 3.2-3.3 (1H, m), 3.4-3.5 (1H, m), 3.5-3.6 (4H, m), 3.7 (1H, t), 3.8 (1H, dd), 4.0 (2H, d), 4.3 (2H, t), 4.8 (1H, dt), 5.0 (1H, d), 5.5 (1H, d), 6.18 (2H, s), 6.7 (2H, d), 7.0-7.1 (3H, m), 7.2 (1H, d), 7.22 (1H, s),
25 7.29 (1H, d), 7.4 (2H, d), 7.8 (2H, d); MS: 791 (MH⁺)

Step 2:

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl) (isobutyl) amino]-2-
 hydroxy-1-{4-[2-(1,3-thiazolidin-3-yl)ethoxy]benzyl}
 propylcarbamate (324)



5

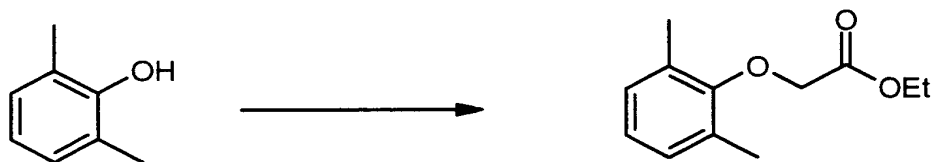
A mixture of 2-(4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-
 Hexahydrofuro[2,3-*b*]furan-3-yloxy]carbonyl}amino)-4-
 [(1,3-benzodioxol-5-ylsulfonyl) (isobutyl) amino]-3-
 10 hydroxybutyl}phenoxy)ethyl 4-methylbenzenesulfonate (50
 mg, 0.06 mmol), thiazolidine (0.02 mL, 0.02 g, 0.25 mmol)
 and dimethylsulfoxide (1 mL) was heated to 70 °C under
 nitrogen atmosphere for 2 h. The mixture was diluted with
 water and extracted with ethyl ether (2X). The organic
 15 phase was dried (sodium sulfate), evaporated, and the
 residue was purified by chromatography (silica gel,
 dichloromethane/methanol, 98:2) to provide the title
 compound (25 mg) as a solid foam. ¹H NMR (DMSO-*d*₆): δ 0.78
 (3H, d), 0.82 (3H, d), 1.2 (1H, dd), 1.4 (1H,
 20 quintuplet), 1.9-2.0 (1H, m), 2.3 (1H, dd), 2.6-2.8 (7H,
 m), 2.8-3.0 (5H, m), 3.4-3.6 (4H, m), 3.7 (1H, t), 3.8
 (1H, dd), 4.0 (2H, br s), 4.05 (2H, br s), 4.8 (1H, dt),
 5.0 (1H, d), 5.5 (1H, d), 6.18 (2H, s), 6.8 (2H, d), 7.0-
 7.1 (3H, d), 7.18 (1H, d), 7.2 (1H, s), 7.3 (1H, d); MS:
 25 708 (MH⁺)

Example 108

Step 1:

Ethyl 2',6'-dimethoxyphenylacetate

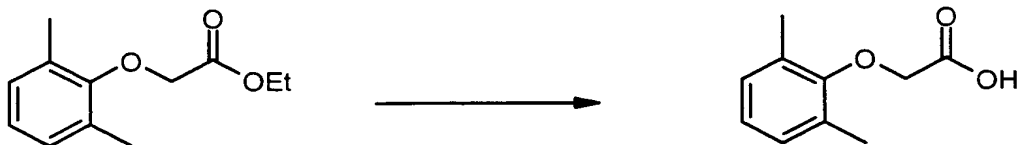
5



- 10 A mixture of 2,6-dimethylphenol (5.18 g, 42.4 mmol),
potassium carbonate (7.33 g, 53.0 mmol) and ethyl
bromoacetate (5.17 mL, 46.6 mmol) in acetone (40 mL) was
stirred at ambient temperature for 24 hours. The reaction
was filtered and the filtered solid was rinsed with
15 acetone. The filtrate was concentrated *in vacuo*, taken up
in ethyl acetate (100 mL), washed with 0.1N sodium
hydroxide (3 x 60 mL), dried (magnesium sulfate) and
concentrated *in vacuo* to afford the crude title compound
(8.79 g, quant.) as a pale yellow liquid. ¹H NMR (DMSO-
20 d₆): δ 1.17 (3H, t), 2.18 (6H, s), 4.14 (2H, q), 4.40 (2H,
s), 6.86-6.98 (3H, m)

Step 2:

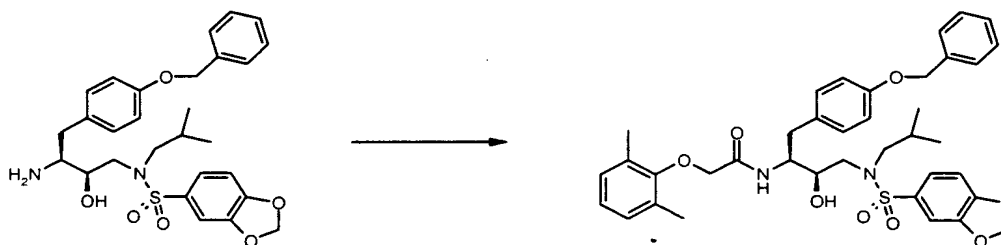
2',6'-Dimethoxyphenylacetic acid



To an ice cold solution of ethyl 2',6'-
dimethylphenoxyacetate (8.79 g, 42.4 mmol) in
tetrahydrofuran/water (120 mL, 3:1) was added lithium
hydroxide monohydrate (3.55 g, 84.8 mmol). After stirring
5 for 3 hours at 5 °C, the tetrahydrofuran was removed in
vacuo and the residual aqueous was diluted with water (70
mL) and extracted with ether (60 mL). The aqueous was
then cooled in an ice bath and acidified to -pH 2 with 1N
hydrochloric acid. The resulting precipitate was
10 extracted into ethyl acetate (150 mL), washed with water
(50 mL), dried (magnesium sulfate) and concentrated in
vacuo to afford the title compound (6.76g, 88%) as a
white solid. ¹H NMR (CDCl₃) : δ 2.28 (6H,s), 4.46 (2H,
s), 6.93-7.01 (3H, m), 10.25 (1H, broad)

Step 3:

N-{(1*S*,2*R*)-3-[(1,3-Benzodioxol-5-ylsulfonyl)(isobutyl)
amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropyl}-2-(2,6-
20 dimethylphenoxy)acetamide (325)

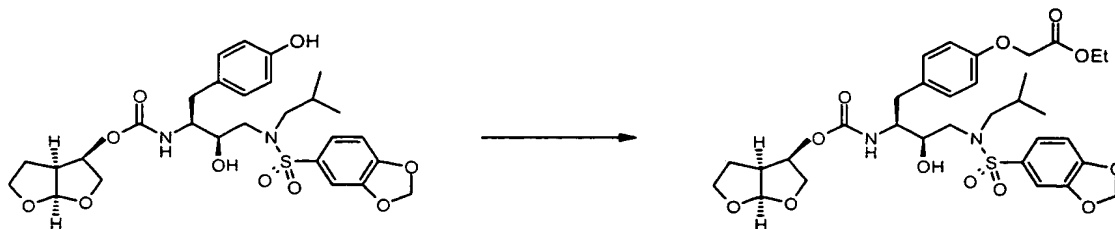


To a stirred solution of *N*-{(2*R*,3*S*)-3-amino-4-[4-
(benzyloxy)phenyl]-2-hydroxybutyl}-*N*-isobutyl-1,3-
benzodioxole-5-sulfonamide (360 mg, 0.68 mmol), 2',6'-
25 dimethylphenoxyacetic acid (160 mg, 0.89 mmol) and *N,N*-
diisopropylethylamine (0.47 mL, 2.7 mmol) in acetonitrile
(7 mL) was added

O-benzotriazol-1-yl-N,N,N',N'-tetramethyluronium
hexafluorophosphate (390 mg, 1.0 mmol). After stirring at
ambient temperature for 1.5 h, the reaction was
concentrated *in vacuo*, taken up in ethyl acetate (60 mL),
5 washed with 0.1N sodium hydroxide (3 x 50 mL) followed by
brine (40 mL), dried (magnesium sulfate) and
concentrated. The residue was purified by silica gel
chromatography (60:40; hexane:ethyl acetate) to afford
the title compound (450 mg, 95%) as a white foam. ¹H NMR
10 (DMSO-d₆): δ 0.78 (6H, dd), 1.94 (1H, m), 2.09 (6H, s),
2.63 (1H, dd), 2.72 (1H, dd), 2.83 (1H, dd), 2.93-3.02
(2H, m), 3.28-3.35 (1H, m), 3.64-3.72 (1H, m), 3.88 (1H,
d), 3.89-3.96 (1H, m), 4.06 (1H, d), 5.01 (2H, s), 5.07
(1H, d), 6.11 (2H, s), 6.82-7.12 (8H, m), 7.25-7.40 (7H,
15 m), 7.86 (1H, d); MS: 689 (MH⁺); C₃₈H₄₄N₂O₈S.

Example 109

20 Ethyl (4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-
b]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-
ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenoxy)
acetate (326)



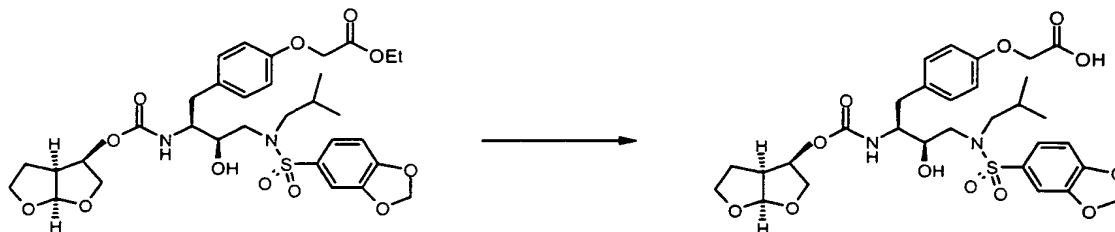
25

To a stirred mixture of (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-
b]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-

ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-hydroxybenzyl)propylcarbamate (200 mg, .34 mmol) and cesium carbonate (330 mg, 1.0 mmol) in N,N-dimethylformamide (5 mL) was added ethyl bromoacetate (75
5 μ L, 0.67 mmol). After stirring at ambient temperature for 2 h, the reaction was diluted with ethyl acetate (50 mL), washed with water (3x40 mL), dried (magnesium sulfate) and concentrated *in vacuo*. The residue was purified by
10 silica gel chromatography (60:40; ethyl acetate:hexane) to afford the title compound (162mg, 71%) as a white foam. ^1H NMR (DMSO- d_6): δ 0.78 (6H, dd), 1.16 (3H, t), 1.17-1.43 (2H, m), 1.92 (1H, m), 2.35 (1H, t), 2.64-2.78 (3H, m), 2.88-3.01 (2H, m), 3.24-3.29 (1H, m), 3.40-3.60 (4H, m), 3.70 (1H, t), 3.78-3.82 (1H, m), 4.10 (2H, q),
15 4.65 (2H, s), 4.81 (1H, q), 5.00 (1H, d), 5.46 (1H, d), 6.12 (2H, s), 6.73 (2H, d), 7.02-7.07 (3H, m), 7.20-7.28 (3H, m); MS: 679 (MH^+)

Example 110

20 (4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenoxy)acetic
25 acid (327)



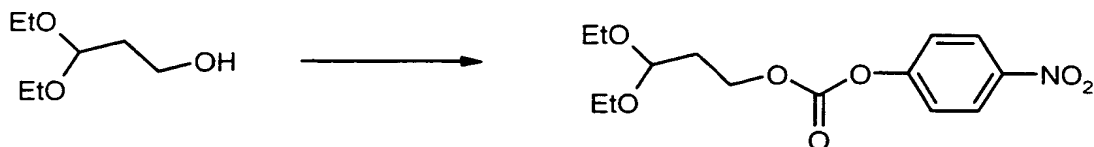
To an ice cold solution of Ethyl (4-{(2*S*,3*R*)-2-
({[(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-
yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-
ylsulfonyl)(isobutyl)amino]-3-
5 hydroxybutyl}phenoxy)acetate (107 mg, 0.16 mmol) in
tetrahydrofuran/water (4 mL, 3:1) was added lithium
hydroxide monohydrate (21 mg, 0.48 mmol). The reaction
was allowed to warm to ambient temperature and stir for 1
h. The tetrahydrofuran was evaporated and the residue
10 diluted with water (30 mL). The aqueous was then
extracted with ether (30 mL), cooled in an ice bath and
acidified to pH ~2 with 1.0 N hydrochloric acid. The
resulting precipitate was extracted into ethyl acetate
(30 mL), washed with water (20 mL), dried (magnesium
15 sulfate) and concentrated *in vacuo* to a white solid.
Triturated with 10% ether in hexane and filtered to
afford the title compound (77 mg) as a 2:1 mixture with
the compound derived from the loss of the bis-
tetrahydrofuran alcohol and subsequent ring closure at
20 the 2-hydroxy position. The mixture was confirmed by NMR,
HPLC and mass spectra. Data for major product: ¹H NMR
(DMSO-*d*₆): δ 0.78 (6H, dd), 1.17-1.45 (2H, m), 1.92 (1H,
m), 2.35 (1H, t), 2.64-2.78 (3H, m), 2.88-3.02 (2H, m),
3.24-3.30 (1H, m), 3.40-3.60 (4H, m), 3.70 (1H, t), 3.78-
25 3.82 (1H, m), 4.54 (2H, s), 4.82 (1H, q), 5.00 (1H, d),
5.46 (1H, d), 6.12 (2H, s), 6.71 (2H, d), 7.00-7.30 (6H,
m), 12.85 (1H, broad); MS: 651 (MH⁺); C₃₀H₃₈N₂O₁₂S (by-
product MS: 521 (MH⁺); C₂₄H₂₈N₂O₉S).

30

Example 328

Step 1:

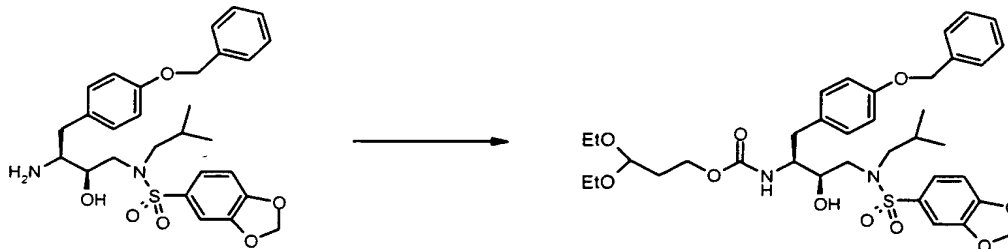
3,3-Diethoxypropan-1-yl-4'-nitrophenyl carbonate



To an ice cold solution of 3,3-diethoxy-1-propanol (1.5 g, 10.1 mmol) and pyridine (1.0 mL, 12.2 mmol) in dichloromethane (30 mL) was added 4-nitrophenylchloroformate (2.24 g, 11.1 mmol). The reaction was allowed to warm to ambient temperature. After stirring for 18 h, the reaction was concentrated in vacuo, taken up in ethyl acetate (60 mL), washed with 5% aqueous citric acid (2x40 mL) followed by saturated sodium carbonate (3x40 mL), dried (magnesium sulfate) and concentrated. The residue was purified by silica gel chromatography (20% ethyl acetate in hexane) to afford the title compound (1.05g, 33%) as an oil. ¹H NMR (CDCl₃): δ 1.19 (6H, t), 2.05 (2H, q), 3.50 (2H, dq), 3.66 (2H, dq), 4.36 (2H, t), 4.66 (1H, t), 7.35 (2H, d), 8.25 (2H, d);

Step 2:

3,3-Diethoxypropyl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate (328)

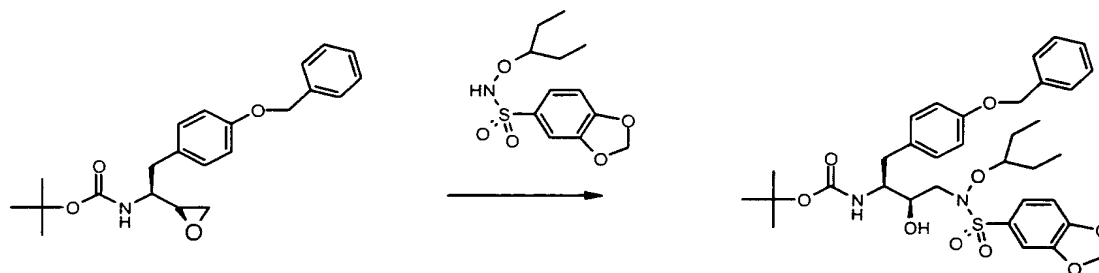


Treatment of *N*-{(2*R*,3*S*)-3-amino-4-[4-(benzyloxy)phenyl]-2-hydroxybutyl}-*N*-isobutyl-1,3-benzodioxole-5-sulfonamide with 3,3-diethoxypropan-1-yl-4'-nitrophenyl carbonate as described above provided the title compound as a white
 5 foam in 51% yield. ¹H NMR (DMSO-d₆): δ 0.78 (6H, dd), 1.33 (6H, dt), 1.65 (2H, q), 1.92 (1H, m), 2.46 (1H, t), 2.69-3.02 (4H, m), 3.27-3.58 (7H, m), 3.80 (2H, t), 4.43 (1H, t), 4.94 (1H, d), 4.99 (2H, s), 6.12 (2H, s), 6.83 (2H, d), 6.97 (1H, d), 7.03 (1H, d), 7.07 (2H, d), 7.23-
 10 7.39 (7H, m); MS: 701 (MH⁺)

Example 112

Step 1:

15 ***tert*-Butyl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(1-ethylpropoxy)amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate**

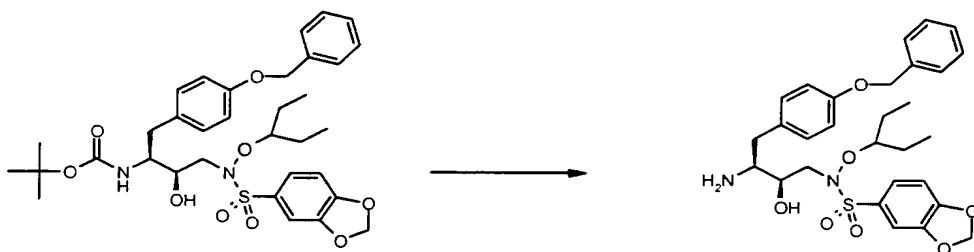


20 To a solution of *tert*-butyl (1*S*)-2-[4-(benzyloxy)phenyl]-1-[(2*S*)-oxiranyl]ethylcarbamate (2.66 g, 7.2 mmol) and the *N*-(1-ethylpropoxy)-1,3-benzodioxole-5-sulfonamide (2.30 g, 9.0 mmol) in tetrahydrofuran (12 mL) was added phosphazene base P4 *tert*-butyl solution (1.44 mL, 1.44
 25 mmol, 1M in hexane). After stirring at ambient temperature for 18 h, the reaction was concentrated in vacuo, taken up in ethyl acetate (100 mL), washed with

0.5 N hydrochloric acid (2x40 mL) followed by water (40 mL), saturated sodium bicarbonate (40 mL) and brine (40 mL), dried (magnesium sulfate) and concentrated. The residue was purified by silica gel chromatography (20% ethyl acetate in hexane) to afford the title compound (4.25 g, 90%) as a white foam. ¹H NMR (DMSO-d₆): δ 0.79 (6H, dt), 1.11-1.74 (4H, m), 1.17 (9H, s), 2.37 (1H, t), 2.60-3.02 (3H, m), 3.38-3.49 (1H, m), 3.50-3.61 (1H, m), 4.03 (1H, m), 5.00 (2H, s), 5.04 (1H, d), 6.15 (2H, s), 6.60 (1H, d), 7.03 (2H, d), 7.10 (1H, d), 7.17-7.38 (9H, m)

Step 2:

N-{(2*R*,3*S*)-3-Amino-4-[4-(benzyloxy)phenyl]-2-hydroxybutyl}-*N*-(1-ethylpropoxy)-1,3-benzodioxole-5-sulfonamide

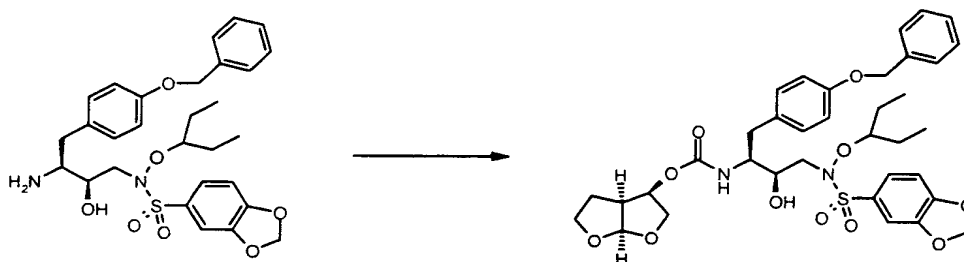


To an ice cold solution of *tert*-Butyl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(1-ethylpropoxy)amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate (1.5 g, 2.3 mmol) in dichloromethane (15 mL) was added trifluoroacetic acid (10 mL). After stirring at 5 °C for 3 h, the reaction was concentrated *in vacuo*, taken up in ethyl acetate (80 mL), washed with saturated sodium bicarbonate (2x50 mL), dried (magnesium sulfate) and concentrated *in vacuo* to afford the title compound (1.27

g, quant.) as an off-white foam. ^1H NMR ($\text{DMSO}-d_6$): δ 0.77 (6H, dt), 1.07-1.72 (4H, m), 1.82 (2H, br), 2.28 (1H, t), 2.58-3.05 (4H, m), 3.43-3.53 (1H, m), 3.95-4.03 (1H, m), 4.86 (1H, br s), 5.01 (2H, s), 6.16 (2H, s), 6.86 (2H, d), 7.03 (2H, d), 7.12-7.40 (8H, m);

Step 3:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(1-ethylpropoxy)amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate (329)

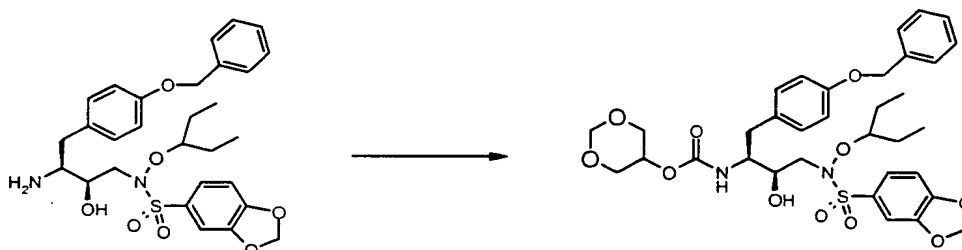


Treatment of *N*-{(2R,3S)-3-Amino-4-[4-(benzyloxy)phenyl]-2-hydroxybutyl}-*N*-(1-ethylpropoxy)-1,3-benzodioxole-5-sulfonamide with [(3R,3aS,6aR)hexahydrofuro[2,3-*b*]furan-3-yl][4-nitrophenyl] carbonate as described above provided the title compound as a white foam in 66% yield. ^1H NMR ($\text{DMSO}-d_6$): δ 0.82 (6H, t), 1.11-1.74 (6H, m), 2.32 (1H, t), 2.69-2.94 (4H, m), 3.45-3.66 (5H, m), 3.75-3.79 (1H, m), 3.99-4.03 (1H, m), 4.78 (1H, q), 5.00 (2H, s), 5.16 (1H, d), 5.46 (1H, d), 6.16 (2H, s), 6.82 (2H, d), 7.03 (2H, d), 7.11-7.38 (9H, m); MS: 713 (MH^+);

$\text{C}_{36}\text{H}_{44}\text{N}_2\text{O}_{11}\text{S}$.

Example 113

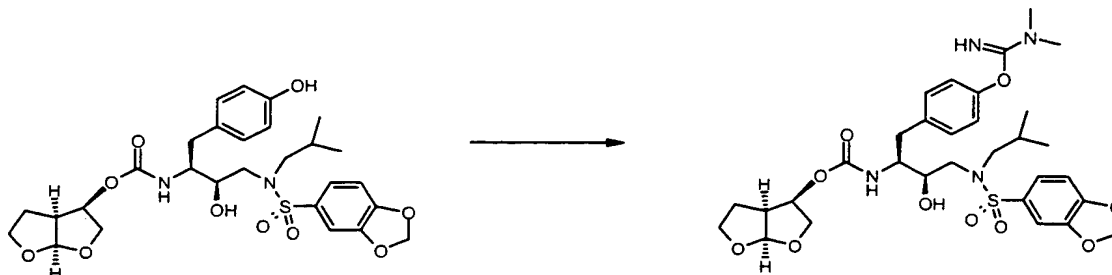
1,3-Dioxan-5-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl) (1-ethylpropoxy) amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate (330)



5 Treatment of *N*-[(2*R*,3*S*)-3-Amino-4-[4-(benzyloxy)phenyl]-2-hydroxybutyl]-*N*-(1-ethylpropoxy)-1,3-benzodioxole-5-sulfonamide with 1,3-dioxan-5-yl-4'-nitrophenyl carbonate as described earlier provided the title compound as a white foam in 65% yield. ¹H NMR (DMSO-*d*₆): δ 0.79 (6H, dt), 1.11-1.74 (4H, m), 2.39 (1H, t), 2.60-3.00 (3H, m), 3.45-3.63 (4H, m), 3.76-3.85 (2H, m), 4.02-4.04 (1H, m), 4.25 (1H, br s), 4.66 (1H, d), 4.74 (1H, d), 4.99 (2H, s), 5.09 (1H, d), 6.16 (2H, s), 6.82 (2H, d), 7.04 (2H, d), 7.12 (1H, d), 7.26-7.39 (7H, m); MS: 687 (MH⁺)

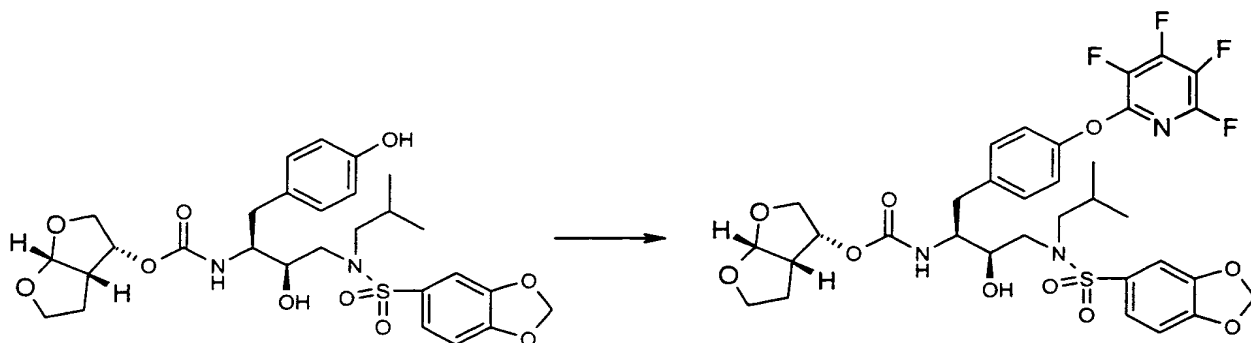
Example 114

(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl) (isobutyl) amino]-1-{4-[(dimethylamino) (imino) methoxy]benzyl}-2-hydroxypropylcarbamate (331)



To an ice cold solution of (3*R*,3*aS*,6*aR*)-
hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-
benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-
hydroxybenzyl)propylcarbamate (100 mg, 0.17 mmol) and
5 cyanogen bromide (36 mg, 0.34 mmol) in acetone (2 mL) was
added a solution of triethylamine (30 μ L, 0.21 mmol) in
acetone (1 mL) over 1 h. After stirring at 5 $^{\circ}$ C for an
additional 2 h, the reaction was concentrated *in vacuo*,
taken up in ethyl acetate (40 mL), washed with water
10 (2x25 mL) followed by brine (25 mL), dried (magnesium
sulfate) and concentrated to afford the crude cyanate
(100 mg) as an oil. The cyanate was dissolved in fresh
acetone (3 mL) and cooled to 5 $^{\circ}$ C. Dimethylamine (3 drops,
2 M in tetrahydrofuran) was added and the reaction was
15 stirred for 30 minutes. The acetone was evaporated and
the residue was purified by silica gel chromatography
(90:10:2; chloroform:methanol:ammonium hydroxide) to
afford the title compound (45 mg, 40%) as a white solid.
 ^1H NMR (DMSO- d_6): δ 0.78 (6H, dd), 1.18-1.45 (2H, m), 1.92
20 (1H, m), 2.44 (1H, t), 2.65-2.78 (3H, m), 2.86-3.02 (3H,
m), 2.88 (6H, s), 3.24-3.29 (1H, m), 3.47-3.61 (4H, m),
3.69 (1H, t), 3.77-3.82 (1H, m), 4.82 (1H, q), 5.05 (1H,
d), 5.47 (1H, d), 6.12 (2H, s), 6.95 (2H, d), 7.03 (1H,
d), 7.17-7.31 (5H, m); MS: 663 (MH $^+$)

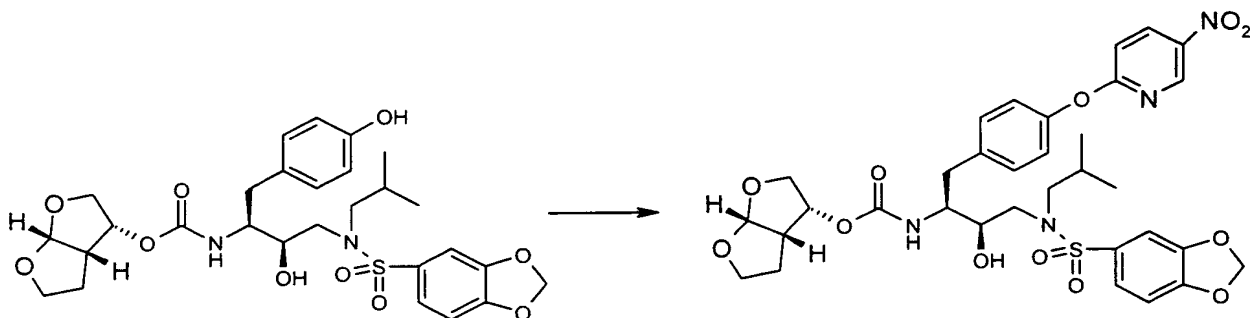
Example 115



5 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-{4-[(3,4,5,6-tetrafluoro-2-pyridinyl)oxy]
benzyl}propylcarbamate (332)

A mixture of 59mg (0.1 mmol) of (3R,3aS,6aR)-
10 hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-
benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-
hydroxybenzyl)propylcarbamate, 51 mg (0.3 mMol) of
pentafluoropyridine and 65mg (0.2 mMol) of cesium
15 carbonate in 0.5 mL of dimethyl formamide was stirred at
rt for 3h. The mixture was diluted with ethyl acetate
and extracted with water. Evaporation of the solvent and
chromatography on silica gel (1:1 ethyl acetate/ hexane)
gave the title compound (32mg) as a white foam. ¹HNMR:
0.85 (6H,dd), 1.55(1H,m), 1.65(1H,m), 1.8(1H,m),
20 2.78(2H,m), 2.9-3.2(5H,m), 3.7(3H,m), 3.8(3H,m),
3.95(1H,m), 4.95(2H,m), 5.62(1H,d), 6.04(2H,s),
6.85(1H,d), 6.95(2H,d), 7.17(1H,s), 7.2(2H,d),
7.34(1H,d). MS: 742 (M+H)

Example 116



5

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-{4-[(5-nitro-2-pyridinyl)oxy]benzyl}
 propylcarbamate (333)

10

A mixture of 59mg (0.1 mmol) of (3*R*,3*aS*,6*aR*)-
 hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-
 benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-
 hydroxybenzyl)propylcarbamate, 42 mg (0.3 mmol) of 2-
 chloro-5-nitropyridine and 65mg (0.2 mmol) of cesium
 carbonate in 0.5 mL of dimethyl formamide was stirred at
 rt for 12h. The mixture was diluted with ethyl acetate
 and extracted with water. Evaporation of the solvent and
 chromatography on silica gel (1:1 ethyl acetate/ hexane)
 gave the title compound (38mg) as a white foam.

15

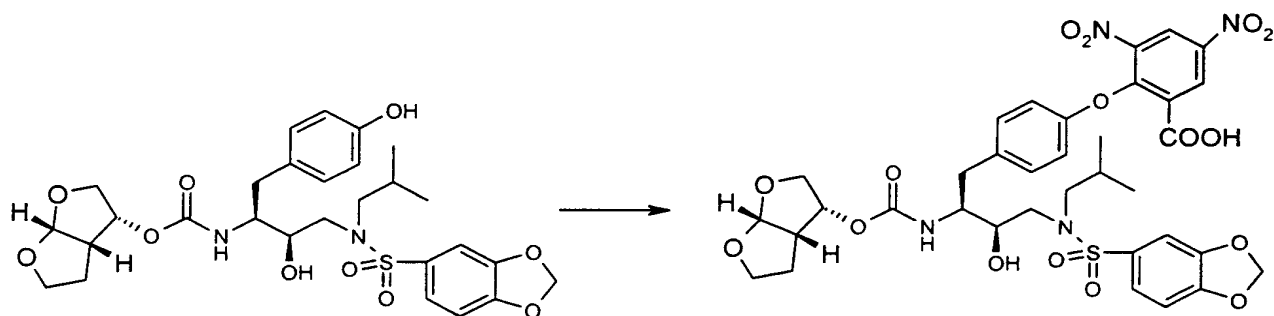
20

¹H-NMR: 0.83 (6H, dd), 1.7 (2H, m), 1.8 (2H, m), 2.8-3.2 (7H, m),
 3.7 (3H, m), 3.8-4 (4H, m), 5.0 (2H, m), 5.62 (1H, d),
 6.03 (2H, s), 6.85 (1H, d), 6.97 (1H, d), 7.02 (2H, d),
 7.18 (1h, s), 7.35 (2H, d), 7.4 (1H, d), 8.42 (1H, d),
 8.90 (1H, d). MS: 715

25

000000-1346500

Example 117

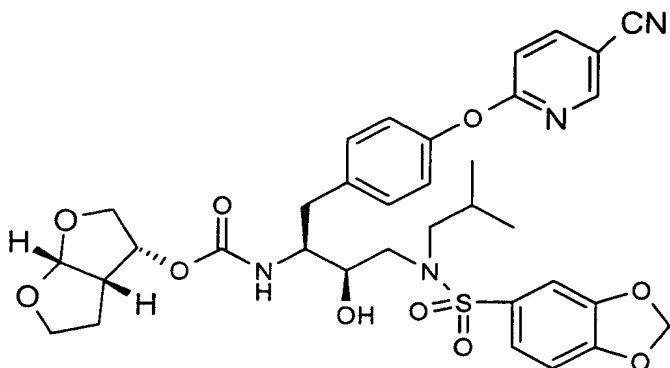


2- (4- { (2*S*,3*R*) -2- ({ [(3*R*,3*aS*,6*aR*) -hexahydrofuro [2,3-
b] furan-3-yloxy] carbonyl} amino) -4- [(1,3-benzodioxol-5-
ylsulfonyl) (isobutyl) amino] -3-hydroxybutyl} phenoxy) -3,5-
dinitrobenzoic acid (334)

A mixture of 59mg (0.1 mMol) of (3*R*,3*aS*,6*aR*) -
hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*) -3- [(1,3-
benzodioxol-5-ylsulfonyl) (isobutyl) amino] -2-hydroxy-1-(4-
hydroxybenzyl)propylcarbamate, 50 of 2-fluoro-3,5-
dinitrobenzoic acid and 65mg (0.2 mMol) of cesium
carbonate in 0.5 mL of dimethyl formamide was stirred at
rt for 2h. The mixture was diluted with ethyl acetate
and extracted with 1N HCl. Evaporation of the solvent
and chromatography on silica gel (5% methanolic ammonia
in dichloromethane) gave the title compound (25mg) as a
yellow foam. ¹H NMR: 0.82(6H,dd), 1.42(1H,m), 1.63(1H,m),
1.90(2H,m), 2.46(1H,dd), 2.8(1H,dd), 2.82-3.02(5H,m),
3.2(1H,m), 3.6-3.95(6H,m), 4.94(1H,q), 5.58(1H,d),
6.02(2H,s), 6.61(1H,d), 6.75(2H,d), 6.82(1H,d),
7.05(2H,d), 7.3(1H,d), 7.45(1H,s), 8.6(2H,ss).

Example 118

5



(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[(5-
 10 cyano-2-pyridinyl)oxy]benzyl}-2-hydroxypropylcarbamate
 (335)

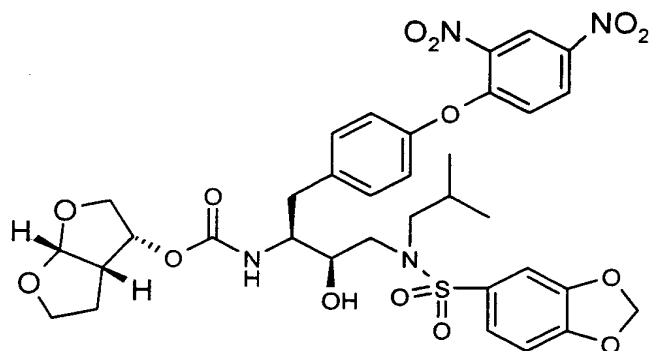
A mixture of 59mg (0.1 mMol) of (3R,3aS,6aR)-
 hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-
 benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-
 15 hydroxybenzyl)propylcarbamate, 120mg of 2-chloro-5-
 cyanopyridine and 65mg (0.2 mMol) of cesium carbonate in
 0.5 mL of dimethyl formamide was stirred at rt for 5h.
 The mixture was diluted with ethyl acetate and extracted
 with water. Evaporation of the solvent and

20 chromatography on silica gel (1:1 ethyl acetate- hexanes)
 gave the title compound (16mg). ¹H NMR: 0.88(6H,dd),
 1.55-1.8(3H,m), 2.8-3.2(7h,m), 3.65(2H,m), 3.8(2H,m),
 3.95(1H,m), 5.0(2H,m), 5.6(1H,d), 6.05(2H,s), 6.82(1H,d),

6.95 (1H,d), 7.0 (2H,d), 7.1-7.4 (4H,m), 7.9 (1H,d),
8.4 (1H,bs). MS: 695 (M+H)

Example 119

5



(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
10 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-
(2,4-dinitrophenoxy)benzyl]-2-hydroxypropylcarbamate
(336)

A mixture of 100mg (0.17 mMol) of (3R,3aS,6aR)-
hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-
15 benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-
hydroxybenzyl)propylcarbamate, 24 mg of 2-chloro-5-
cyanopyridine and 27mg of triethylamine in 0.5 mL of
dimethyl formamide was stirred at rt for 5h. The mixture
was diluted with ethyl acetate and extracted with 1N HCl
20 and water. Evaporation of the solvent and chromatography
on silica gel (1:1 ethyl acetate- hexanes) gave the title
compound (140mg) as a yellow solid. ¹H NMR: 0.9(6H,dd),
1.6-1.8(3H,m), 2.8-3.2(9H,m), 3.7(3H,m), 3.8-4(4H,m),
5.0(2H,m), 5.6(1H,d), 6.05(2H,s), 6.85(1H,d), 6.99(1H,d),

7.05 (2H,d), 7.25 (1H,s), 7.3 (3H,m), 8.3 (1H,dd), 9.0 (1H,d).

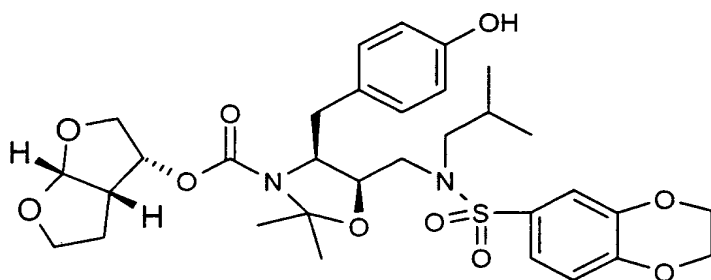
MS: 759 (M+H)

Example 120

5

Step 1:

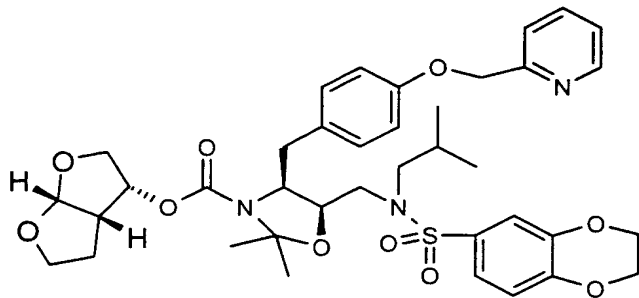
(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[(2,3-dihydro-1,4-benzodioxin-6-
10 ylsulfonyl)(isobutyl)amino]methyl}-4-(4-hydroxybenzyl)-
2,2-dimethyl-1,3-oxazolidine-3-carboxylate



15 250mg of (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl
(1*S*,2*R*)-3-[(2,3-dihydro-1,4-benzodioxin-6-
ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-
benzyloxybenzyl)propylcarbamate, 2mL of 2,2-
dimethoxypropane, 0.5g of p-toluenesulfonic acid in 50mL
20 of dichloromethane were refluxed for 3h and then
extracted with sodium bicarbonate solution.
Chromatography on silica gel (1:1 ethylacetate/hexanes)
gave 220mg of the desired compound as an oil which was
dissolved in methanol and hydrogenated (at 50 psi) over
25 5% palladium on carbon. Filtration and evaporation of
the volatiles gave the desired title compound as an oil

Step 2:

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (4S,5R)-5-
 {[(2,3-dihydro-1,4-benzodioxin-6-
 5 ylsulfonyl)(isobutyl)amino]methyl}-2,2-dimethyl-4-[4-(2-
 pyridinylmethoxy)benzyl]-1,3-oxazolidine-3-carboxylate

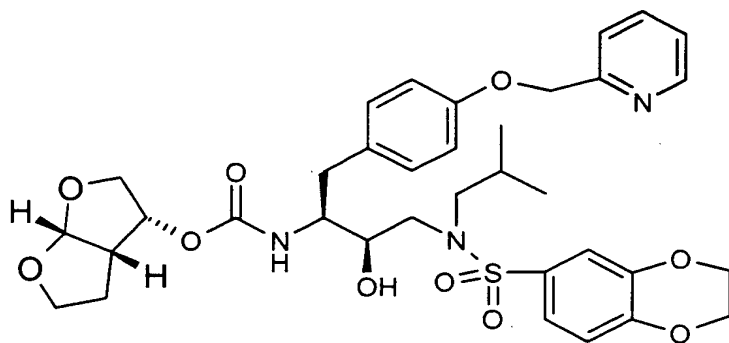


0.1g of (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl
 10 (4S,5R)-5-{[(2,3-dihydro-1,4-benzodioxin-6-
 ylsulfonyl)(isobutyl)amino]methyl}-4-(4-hydroxybenzyl)-
 2,2-dimethyl-1,3-oxazolidine-3-carboxylate, 0.03g of o-
 picolylchloride-hydrochloride and 0.15g of cesium
 carbonate were suspended in 0.5 mL of dimethyl formamide
 15 and heated to 60 degrees for 3h. The mixture was diluted
 with ethyl acetate and washed with water. Chromatography
 on silica gel (9:1 ethyl acetate/hexanes) gave 72mg of
 the desired material

20 Step3:

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(2,3-dihydro-1,4-benzodioxin-6-
 ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-[4-(2-
 pyridinylmethoxy)benzyl]propylcarbamate (337)

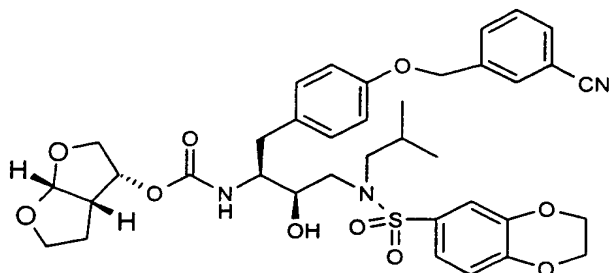
25



70mg of (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl
(4*S*,5*R*)-5-{ [(2,3-dihydro-1,4-benzodioxin-6-
5 ylsulfonyl) (isobutyl) amino] methyl }-2,2-dimethyl-4-[4-(2-
pyridinylmethoxy) benzyl]-1,3-oxazolidine-3-carboxylate
were dissolved in 15mL of isopropanol and 5mL of
concentrated hydrochloric acid. After 2h, the reaction
was treated with excess 3N sodium hydroxide and extracted
10 with ethyl acetate. Evaporation of the solvent gave 67mg
of the desired product. ¹H NMR: 0.83(6H,dd), 1.4-
1.8(4H,m), 2.6-3.2(7H,m), 3.6(2H,m), 3.75(2H,m),
3.9(2H,m), 4.25(4H,bs), 4.95(1H,d), 5.0(1H,m), 5.6(1H,d),
6.9(2H,d), 6.95(2H,d), 7.1(2H,d), 7.2(2H,m), 7.45(1H,d),
15 7.7(1H,t), 8.58(1H,d).

Example 121

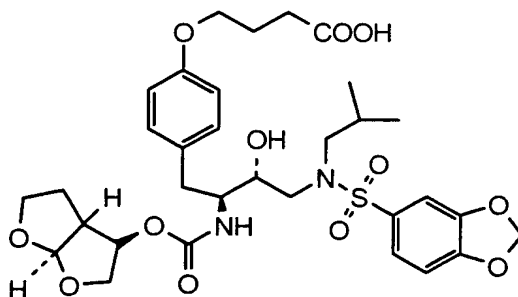
(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-{4-
20 [(3-cyanobenzyl) oxy] benzyl }-3-[(2,3-dihydro-1,4-
benzodioxin-6-ylsulfonyl) (isobutyl) amino]-2-
hydroxypropylcarbamate (338)



49mg of (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl
 (1*S*,2*R*)-3-[(2,3-dihydro-1,4-benzodioxin-6-
 ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-hydroxy
 5 benzyl)propylcarbamate, 20mg of *m*-cyanobenzylchloride,
 20mg of cesium carbonate and 0.5 mL of dimethyl formamide
 were stirred at rt for 5h. The mixture was diluted with
 ethyl acetate and extracted with water. Chromatography
 on silica gel (1:1 ethylacetate/hexane) gave the title
 10 compound as a white solid (26mg). ¹H NMR: 0.87(6H,dd),
 1.58(2H,m), 1.8(1H,m), 2.75(2H,m), 2.9(4H,m), 3.1(1H,m),
 3.62(3H,m), 3.8(3H,m), 3.95(1H,m), 4.25(4H,m),
 4.95(2H,m), 5.0(2H,s), 5.62(1H,d), 6.82(2H,d),
 6.92(1H,d), 7.1(2H,d), 7.2(2H,m), 7.45(1H,t), 7.6(2H,m),
 15 7.75(1H,s). MS:722 (M+H)

Example 122

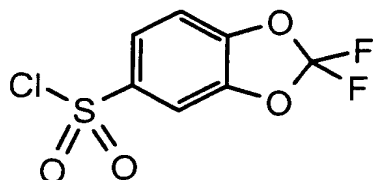
4-(4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-
b]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-
 20 ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenoxy)
 butanoic acid (339)



To a 0° C solution of Methyl 4-(4-{(2S,3R)-2-
({[3R,3aS,6aR]-hexahydrofuro[2,3-b]furan-3-
yloxy}carbonyl)amino)-4-[(1,3-benzodioxol-5-
5. ylsulfonyl)(isobutyl)amino]-3-
hydroxybutyl}phenoxy)butanoate (0.9 g, 0.144 mmol) in 8
ml of THF-H₂O (3:1) was added lithium hydroxide
monohydrate (30 mg, 0.715 mmol). The ice bath was removed
and the reaction mixture was stirred at room temperature
10 for 2 hours. Carefully acidified to pH 2 with 1N HCl and
diluted with 50 ml of dichloromethane. The organic layer
was washed with water and then dried with sodium sulfate,
filtered and concentrated under reduced pressure.
Residue was triturated with diethyl ether and filtered to
15 afford 85 mg (87%) of the title compound.
¹H-NMR: (DMSO-d₆): δ 0.79 (3H,d), 0.83 (3H,d), 1.20-1.31
(2H,m), 1.35-1.45 (2H,m) 1.80-1.98 (3H,m), 2.30-2.41
(2H,m), 2.65-2.82 (3H,m), 2.85-3.06 (3H,m), 3.34-3.61
(4H,m), 3.70-3.76 (1H,m), 3.82-3.98 (3H,m), 4.84 (1H,q),
20 5.01 (1H,d), 5.51 (1H,d), 6.17 (2H,s), 6.77 (2H,d), 6.88
(1H,d), 7.01-7.18 (3H,m), 7.23 (1H,d), 12.06 (1H,br s).
MS: 679 (M⁺). C₃₂H₄₂N₂O₁₂S.

Example 123

25 Step 1:
2,2-difluoro-1,3-benzodioxole-5-sulfonyl chloride

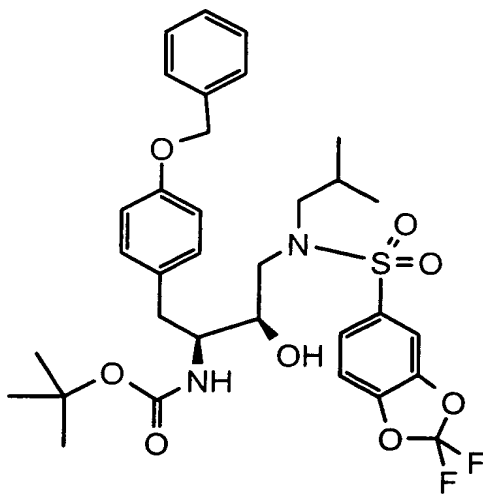


A solution of 2.37 g (10 mmol) 5-bromo-2,2-difluoro-1,3-benzodioxole in 25 ml anhydrous ether, cooled to -30 °C, and 4 ml (10 mmol) *n* butyllithium (2.5 M in hexanes) was added at -30 °C, over 10 minutes. The mixture was stirred at -30 °C for 1 hour, then sulfur dioxide gas was passed through the mixture at -20 °C for 5 minutes. The mixture then was allowed to warm up to 20 °C, and the precipitate was filtered and washed with 20 ml hexane. The solid was suspended in 20 ml hexanes, 1.35 g (10 mmol) sulfuryl chloride was added, the mixture was stirred at 20 °C for 48 hours. The mixture then was filtered and the filtrate was concentrated under reduced pressure to obtain 1.6 g (62 %) title compound. The product was used in the next step without further purification.

¹H-NMR (CDCl₃): δ 7.27 (1H,d), 7.74 (1H,s), 7.88 (1H,d).

Step 2:

tert-butyl (1*S*,2*R*)-1-[4-(benzyloxy)benzyl]-3-[[2,2-difluoro-1,3-benzodioxol-5-yl)sulfonyl](isobutyl)amino]-2-hydroxypropylcarbamate



The sulfonamide formation was carried out as described for t-Butyl-N((1S,2R)-1-(4-benzyloxy-benzyl)-3-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carbamate. (Y=73%)

5

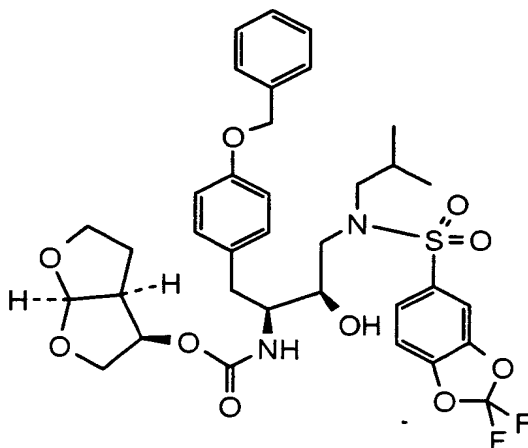
¹H-NMR (CDCl₃): δ 0.86 (3H,d), 0.88 (3H,d), 1.34 (9H,s), 1.84 (1H,m), 2.85 (4H,m), 3.09 (2H,d), 3.68 (1H,s(br)), 3.78 (2H,m), 4.57 (1H,d(br)), 5.02 (2H,s), 6.90 (2H,d), 7.13 (2H,d), 7.15-60 (8H,m).

10

Step 3:

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-1-[4-(benzyloxy)benzyl]-3-[[[(2,2-difluoro-1,3-benzodioxol-5-yl)sulfonyl](isobutyl)amino]-2-hydroxypropyl]carbamate (340)

15



The carbamate formation was carried out as described for (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-benzyloxy-benzyl)-3-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carbamate. (Y=48%)

20

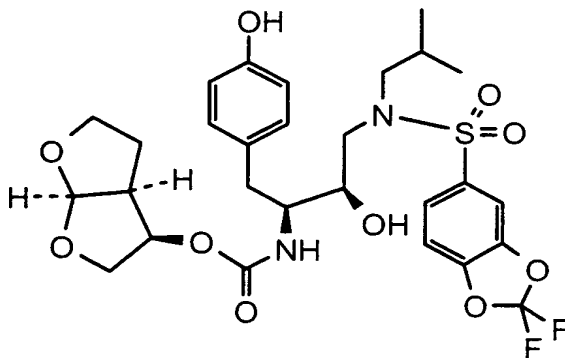
¹H-NMR (DMSO-d₆): δ 0.77 (3H,d), 0.82 (3H,d), 1.20 (1H,m),
1.35 (1H,m), 1.90 (1H,m), 2.35 (1H,t), 2.70-3.00 (5H,m),
3.04 (1H,dd), 3.40-3.60 (4H,m), 3.66 (1H,t), 3.80
(1H,dd), 4.81 (1H,dd), 4.98 (1H,d), 4.99 (2H,s), 5.47
5 (1H,d), 6.82 (2H,d), 7.06 (2H,d), 7.18 (1H,d), 7.20-7.45
(5H,m), 7.56 (1H,d), 7.65 (1H,d), 7.81 (1H,s). MS: 719
(M+1)⁺.

Example 124

10

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
[[(2,2-difluoro-1,3-benzodioxol-5-
yl) sulfonyl] (isobutyl) amino] -2-hydroxy-1-(4-
hydroxybenzyl)propylcarbamate (341)

15



20 The debenzylation was carried out as described for N-
(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-,
(4S,5R)-4-(4-hydroxybenzyl)-5-i-butyl-[(3,4-
methylenedioxyphenyl) sulfonyl]-aminomethyl-2,2-dimethyl-
oxazolidine. (Y=100%)

25

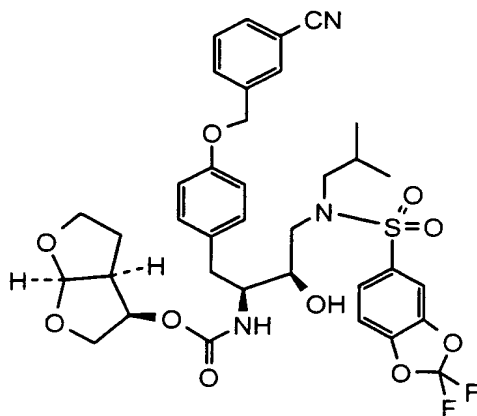
000090" 4946560

¹H-NMR (DMSO-d₆): δ 0.77 (3H,d), 0.82 (3H,d), 1.25 (1H,m),
1.40 (1H,m), 1.92 (1H,m), 2.30 (1H,t), 2.70-3.00 (5H,m),
3.02 (1H,dd), 3.40-3.60 (4H,m), 3.70 (1H,t), 3.80
(1H,dd), 4.82 (1H,dd), 4.95 (1H,d), 5.47 (1H,d), 6.56
5 (2H,d), 6.92 (2H,d), 7.13 (1H,d), 7.56 (1H,d), 7.64
(1H,d), 7.81 (1H,s), 9.00 (1H,s).

10

Example 125

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-1-{4-
[(3-cyanobenzyl)oxy]benzyl}-3-[[[(2,2-difluoro-1,3-
benzodioxol-5-yl)sulfonyl](isobutyl)amino]-2-
15 hydroxypropylcarbamate (342)



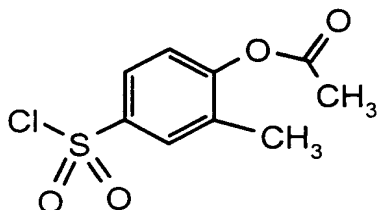
The alkylation step was carried out as described for N-
20 (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-,
(4S,5R)-4-[4-(3-cyanobenzoyloxy)-benzyl]-5-i-butyl-[(3,4-
methylenedioxyphenyl)sulfonyl]-aminomethyl-2,2-dimethyl-
oxazolidine. (Y=84%)

¹H-NMR (DMSO-d₆): δ 0.77 (3H,d), 0.81 (3H,d), 1.21 (1H,m),
1.32 (1H,m), 1.92 (1H,m), 2.35 (1H,t), 2.65-3.00 (5H,m),
3.05 (1H,t), 3.40-3.60 (4H,m), 3.65 (1H,t(br)), 3.80
(1H,t(br)), 4.81 (1H,d(br)), 4.99 (1H,m), 5.08 (2H,s),
5 5.47 (1H,d), 6.84 (2H,d), 7.07 (2H,d), 7.19 (1H,d), 7.50-
7.90 (7H,m). MS: 744 (M+1)⁺.

Example 126

10 Step 1:

4-(chlorosulfonyl)-2-methylphenyl acetate



21 g (205 mmol) acetic anhydride was added to a stirred
15 mixture of 18.8 g (100 mmol) 3-methyl, 4-hydroxy sulfonic
acid in 100 ml dichloromethane. The mixture was stirred
for 12 hours, then the solvent was removed under reduced
pressure. The residue was dissolved in 100 ml ether, and
75 g (364 mmol) phosphorus pentachloride was added to the
20 stirred solution in small portions. After the addition,
the mixture was stirred for 15 minutes at 20 °C, then
poured into 500 g crushed ice. An additional 200 ml ether
was added, and the organic phase was separated, extracted
with cold water, then dried over anhydrous magnesium
25 sulfate. The solvent was removed under reduced pressure,
150 ml ether and 150 ml hexane was added, and the solid
was filtered. The filtrate was concentrated under reduced

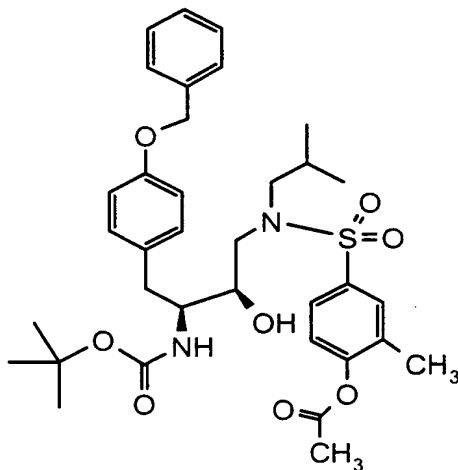
pressure again, then hexane (neat) was added, and the mixture was cooled to 0 °C. The solid was filtered, washed with cold hexane, and dried under reduced pressure, to yield 6.8 g (Y=27%) title compound.

5

¹H-NMR (CDCl₃): δ 2.29 (3H,s), 2.36 (3H,s), 7.27 (1H,d), 7.88 (1H,d), 7.91 (1H,s).

10 Step 2:

4-{[{(2R,3S)-4-[4-(benzyloxy)phenyl]-3-[(tert-butoxycarbonyl)amino]-2-hydroxybutyl}(isobutyl)amino]sulfonyl}-2-methylphenyl
15 acetate



The sulfonamide formation was carried out as described for t-Butyl-N((1S,2R)-1-(4-benzyloxy-benzyl)-3-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carbamate. (Y=74%)

20

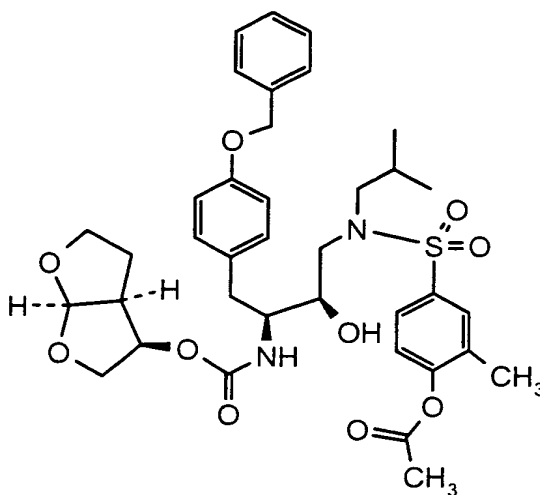
¹H-NMR (CDCl₃): δ 0.85 (3H,d), 0.89 (3H,d), 1.33 (9H,s), 1.84 (1H,m), 2.23 (3H,s), 2.32 (3H,s), 2.80-3.15 (6H,m),

3.86 (1H,s(br)), 3.77 (1H,s(br)), 3.90 (1H, s), 4.59 (1H,d), 5.02 (2H,s), 6.89 (2H,d), 7.10-7.25 (8H,m), 7.59 (1H,d), 7.64 (1H,s).

5 Step 3:

4-{ [{ (2R,3S) -3- ({ [(3R,3aS,6aR) -hexahydrofuro[2,3-b]furan-3-yloxy] carbonyl } amino) -4- [4- (benzyloxy)phenyl] -2-hydroxybutyl} (isobutyl) amino] sulfonyl} -2-methylphenyl acetate (343)

10



The carbamate formation was carried out as described for (3R,3aS,6aR) -Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R) -1-(4-benzyloxy-benzyl) -3-i-butyl- [(3,4-methylenedioxyphenyl) sulfonyl] -amino-2-hydroxypropyl)-carbamate. (Y=29%)

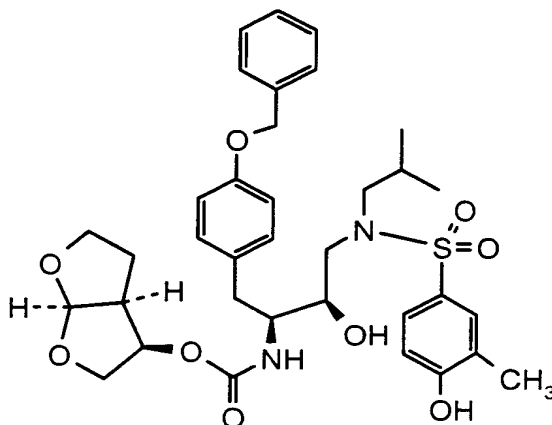
¹H-NMR (DMSO-d₆): δ 0.76 (3H,d), 0.81 (3H,d), 1.21 (2H,m), 1.32 (1H,m), 1.93 (1H,m), 2.16 (3H,s), 2.29 (3H,s), 2.36 (1H,t), 2.70-2.90 (3H,m), 2.93 (1H,d), 3.03 (1H,dd), 3.35 (1H,m), 3.45-3.60 (4H,m), 3.66 (1H, dd), 3.80 (1H,dd),

4.80 (1H,dd), 4.99 (3H,s), 5.46 (1H,d), 6.82 (2H,d), 7.07 (2H,d), 7.18 (1H,d), 7.22-7.42 (5H,m), 7.61 (1H,d), 7.70 (1H,s).

5

Example 127

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-(benzyloxy)benzyl]-2-hydroxy-3-[[4-hydroxy-3-methylphenyl)sulfonyl](isobutyl)amino]propylcarbamate (344)



10

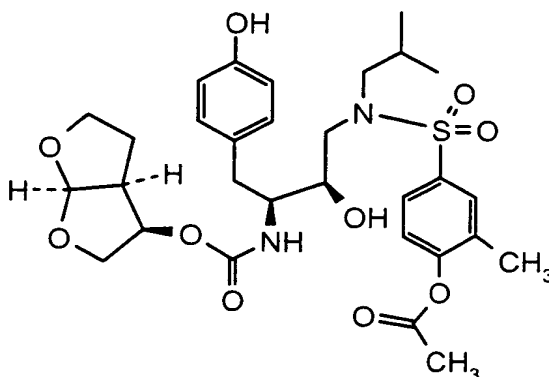
100 mg (0.14 mmol) product from the previous procedure dissolved in 10 ml methanol. 12 mg potassium carbonate was added to the solution, and the mixture was stirred for 30 minutes at 20 °C. The solvent was removed under reduced pressure, and the residue was dissolved in dioxane. The pH was adjusted to pH=4 with hydrochloric acid (4M in dioxane), the mixture was filtered, and the filtrate was concentrated under reduced pressure. The product crystallized from hexane-ethyl acetate (1:1) at -20 °C the yield 75 mg title compound. (Y=79%).

20

¹H-NMR (DMSO-d₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.20 (1H,m), 1.30 (1H,m), 1.90 (1H,m), 2.12 (3H,s), 2.35 (1H,t), 2.60-2.80 (3H,m), 2.85-3.00 (2H,m), 3.40-3.60 (4H,m), 3.66 (1H,dd), 3.81 (1H,dd), 4.80 (1H,dd), 4.94 (1H,d), 5.00 (2H,s), 5.47 (1H,d), 6.85 (3H,m), 7.06 (2H,d), 7.18 (1H,d), 7.30-7.50 (6H,m), 10.3 (1H,s). MS: 669 (M+1)⁺.

Example 128

4-{[[(2R,3S)-3-({[(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yloxy]carbonyl}amino)-2-hydroxy-4-(4-hydroxyphenyl)butyl](isobutyl)amino]sulfonyl}-2-methylphenyl acetate (345)



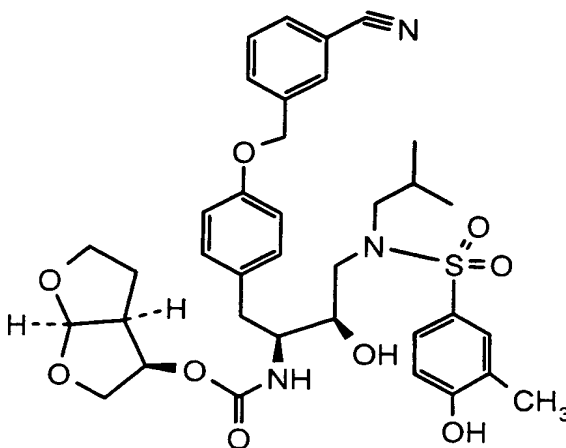
The debenzylation was carried out as described for N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-oxy carbonyl-, (4S,5R)-4-(4-hydroxybenzyl)-5-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-aminomethyl-2,2-dimethyloxazolidine. (Y=quant.)

¹H-NMR (CDCl₃): δ 0.85 (3H,d), 0.88 (3H,d), 1.45 (1H,m), 1.62 (1H,m), 2.21 (3H,s), 2.32 (3H,s), 2.65 (1H,dd), 2.80-3.15 (6H,m), 3.60-3.70 (3H,m), 3.75-3.95 (3H,m),

5.00 (1H,m), 5.21 (1H,d), 5.60 (1H,d), 6.65 (2H,d), 6.90-7.00 (3H,m), 7.15 (1H,d), 7.56 (1H,d), 7.62 (1H,s).

Example 129

5 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-1-{4-[(3-cyanobenzyl)oxy]benzyl}-2-hydroxy-3-[[4-(4-hydroxy-3-methylphenyl)sulfonyl](isobutyl)amino]propylcarbamate (346)



10

The alkylation step was carried out as described for N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-, (4S,5R)-4-[4-(3-cyanobenzyloxy)-benzyl]-5-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-aminomethyl-2,2-dimethyl-oxazolidine. The deacetylation step was carried out as described previously for (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-1-[4-(benzyloxy)benzyl]-2-hydroxy-3-[[4-(4-hydroxy-3-methylphenyl)sulfonyl](isobutyl)amino]propylcarbamate.

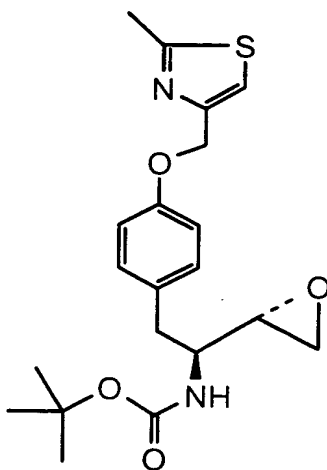
20 The final product was isolated by preparative HPLC. (Column: Luna C18, Solvent: 75% to 100 % MeOH/0.1%formic acid, t_{ret} : 4.08 min.) (Y=6 %, 2 steps, overall.)

MS: 694 (M+1)

Example 130

Step 1:

- 5 ***tert*-butyl (1*S*)-2-{4-[(2-methyl-1,3-thiazol-4-yl) methoxy]phenyl}-1-[(2*S*)-oxiranyl]ethylcarbamate**

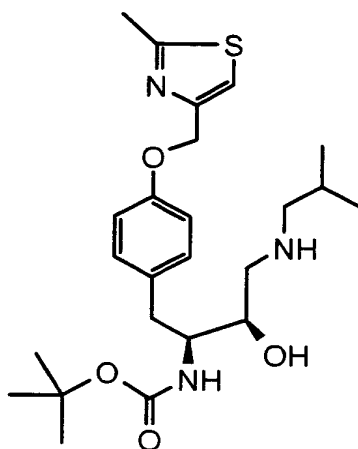


- 10 0.6 g (3.26 mmol) 4-Chloromethyl-2-methyl-thiazole hydrochloride was added to a stirred solution of 0.76 g (2.71 mmol) *tert*-butyl (1*S*)-2-(4-hydroxyphenyl)-1-[(2*S*)-oxiranyl]ethylcarbamate in 15 ml *N,N*-dimethylformamide. 3.25 g (10 mmol) cesium carbonate and 550 mg (3.66 mmol)
- 15 sodium iodide was added, and the mixture was stirred at 20 °C for 24 hours. 200 ml ethyl acetate was added, and the mixture was filtered. The filtrate was extracted with water (5x50 ml), and the organic phase was dried with anhydrous magnesium sulfate, filtered and concentrated
- 20 under reduced pressure. The residue was purified on silicagel column using hexane-ethyl acetate (1:1) as solvent to yield 820 mg (Y=78 %) title compound.

¹H-NMR (CDCl₃): δ 1.36 (9H,s), 2.71 (3H,s), 2.75-2.95 (4H,m), 3.61 (1H,s(br)), 4.39 (1H,s(br)), 5.11 (2H,s), 6.91 (2H,d), 7.11 (2H,d), 7.12 (1H,s).

5 Step 2:

tert-butyl (1S,2R)-2-hydroxy-3-(isobutylamino)-1-{4-[(2-methyl-1,3-thiazol-4-yl)methoxy]benzyl}propylcarbamate



10 The ring-opening step was carried out as described previously for t-Butyl-(1S,2R)-1-(4-benzyloxy-benzyl)-3-i-butylamino-2-hydroxypropyl-carmamate. (Y=90%)

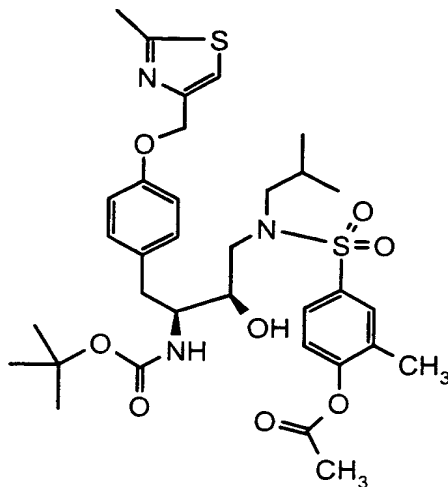
15 ¹H-NMR (CDCl₃): δ 0.94 (6H,d), 1.39 (9H,s), 1.74 (1H,m), 2.43 (2H,d), 2.70 (2H,m), 2.76 (3H,s), 2.80-3.00 (2H,m), 3.46 (1H,m), 3.79 (1H,m), 4.68 (1H,d), 5.16 (2H,s), 6.94 (2H,d), 7.17 (1H,s), 7.18 (2H,d).

Step 3:

20

4-{[((2R,3S)-3-[(tert-butoxycarbonyl)amino]-2-hydroxy-4-{4-[(2-methyl-1,3-thiazol-4-

yl)methoxy]phenyl}butyl) (isobutyl) amino] sulfonyl}-2-methylphenyl acetate



5

The sulfonamide formation was carried out as described for t-Butyl-N((1S,2R)-1-(4-benzyloxy-benzyl)-3-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carbamate. (Y=65%)

10

¹H-NMR (CDCl₃): δ 0.85 (3H,d), 0.87 (3H,d), 1.33 (9H,s), 1.82 (1H,m), 2.23 (3H,s), 2.32 (3H,s), 2.71 (3H, s), 2.80-3.00 (4H,m), 3.10 (2H,m), 3.68 (1H,s(br)), 3.77 (1H,s(br)), 3.91 (1H,s), 4.57 (1H,d), 5.10 (2H,s), 6.90 (2H,d), 7.12 (1H,s), 7.13 (2H,d), 7.59 (1H,d), 7.64 (1H,s).

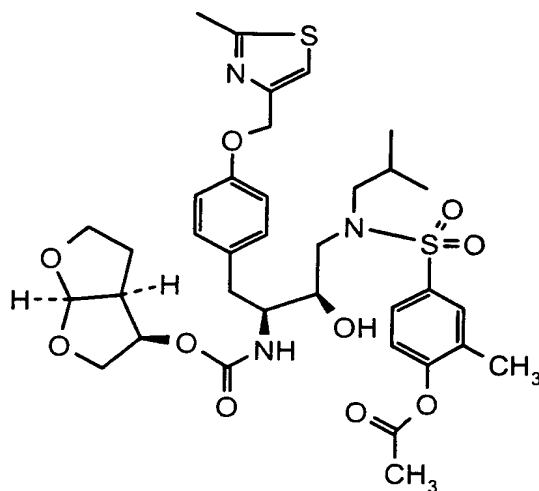
15

Step 4:

20

4-{[(2R,3S)-3-({[(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yloxy]carbonyl}amino)-2-hydroxy-4-{4-[(2-methyl-1,3-

thiazol-4-yl)methoxy]phenyl}butyl) (isobutyl) amino]
sulfonyl}-2-methylphenyl acetate (347)



5 The carbamate formation was carried out as described for
(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-
(4-benzyloxy-benzyl)-3-i-butyl-[(3,4-
methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-
carmamate. (Y=44%)

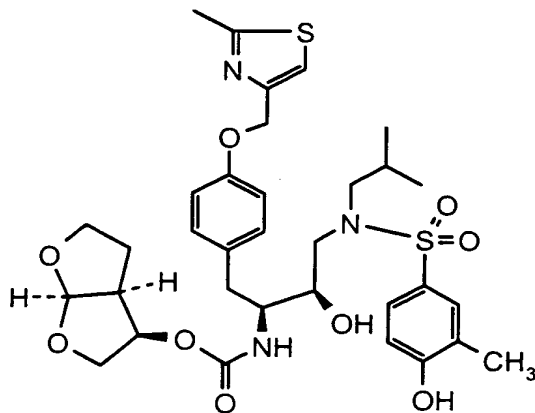
10

¹H-NMR (DMSO-d₆): δ 0.76 (3H,d), 0.81 (3H,d), 1.34 (1H,m),
1.94 (1H,m), 2.16 (3H,s), 2.30 (3H,s), 2.36 (1H,t), 2.61
(3H,s), 2.70-3.10 (5H,m), 3.45-3.60 (3,m), 3.68 (1H,t),
3.79 (1H,t), 4.81 (1H,m), 4.99 (3H, sm), 5.46 (1H,d),
15 6.84 (2H,d), 7.08 (2H,d), 7.20 (1H,d), 7.26 (1H,d), 7.48
(1H,d), 7.61 (1H,d), 7.70 (1H,s). MS: 732 (M+1)⁺.

Example 131

20 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-2-
hydroxy-3-[[[4-hydroxy-3-methylphenyl)sulfonyl](isobutyl)

amino]-1-{4-[(2-methyl-1,3-thiazol-4-yl)methoxy]benzyl}
propylcarbamate (348)



5

Procedure:

The deacetylation step was carried out as described previously for (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-1-[4-(benzyloxy)benzyl]-2-hydroxy-3-[[4-hydroxy-3-methylphenyl)sulfonyl](isobutyl)amino] propylcarbamate. (Y=94%)

10

¹H-NMR (DMSO-*d*₆): δ 0.75 (3H,s(br)), 0.81 (3H,s(br)), 1.20 (1H,m), 1.32 (1H,m), 1.91 (1H,m), 2.12 (3H,s), 2.36 (1H,t), 2.61 (3H,s), 2.65-3.00 (5H,m), 3.40-3.60 (4H,m), 3.68 (1H,s(br)), 3.81 (1H,s(br)), 4.80 (1H,m), 4.99 (3H,s+m), 5.40 (1H,s(br)), 6.85 (4H,m), 7.07 (2H,m), 7.17 (1H,d), 7.35-7.50 (3H,m), 10.30 (1H,s). MS: 690 (M+1)⁺.

15

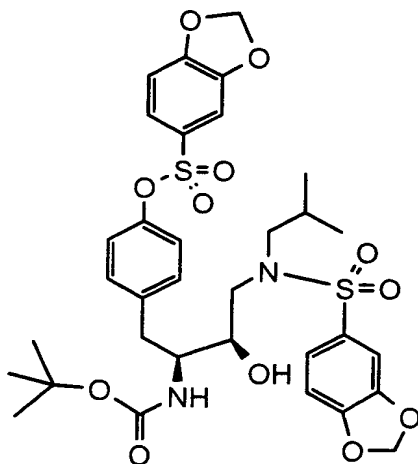
20

Example 132

Step 1:

4-{(2*S*,3*R*)-4-[(1,3-benzodioxol-5-yl)sulfonyl](isobutyl)amino]-2-[(tert-

butoxycarbonyl)amino]-3-hydroxybutyl}phenyl 1,3-benzodioxole-5-sulfonate



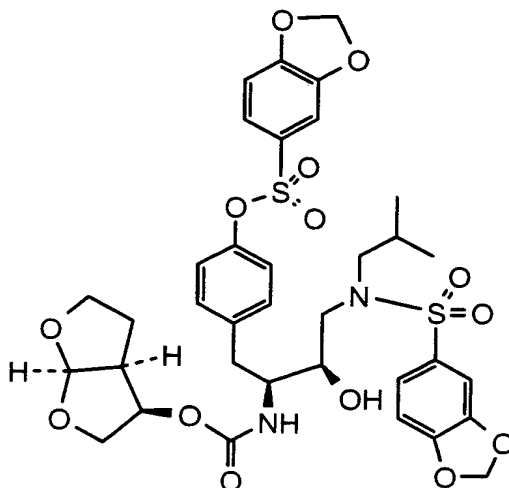
5

The reaction was carried as described for the synthesis of 4-((2*S*,3*R*)-2-[(*tert*-butoxycarbonyl)amino]-3-hydroxy-4-
10 {isobutyl[(4-nitrophenyl)sulfonyl]amino}butyl)phenyl 4-nitrobenzenesulfonate.

¹H-NMR (CDCl₃): δ 0.86 (3H,d), 0.89 (3H,d), 1.33 (9H,s),
1.81 (1H,m), 2.75-3.10 (6H,m), 3.68 (1H,s(br)), 3.76
15 (1H,s(br)), 3.95 (1H,s), 4.60 (1H,d), 6.07 (2H,s), 6.09
(2H,s), 6.80-6.95 (4H, m), 7.10-7.35 (6H,m).

Step 2:

20 4-{(2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yloxy]carbonyl}amino)-4-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-3-hydroxybutyl}phenyl 1,3-benzodioxole-5-sulfonate (349)



5

The reaction was carried as described for the synthesis of 4-((2*S*,3*R*)-2-({[(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yloxy]carbonyl}amino)-3-hydroxy-4-{isobutyl[(4-nitrophenyl)sulfonyl]amino}butyl)phenyl 4-nitrobenzenesulfonate.

10

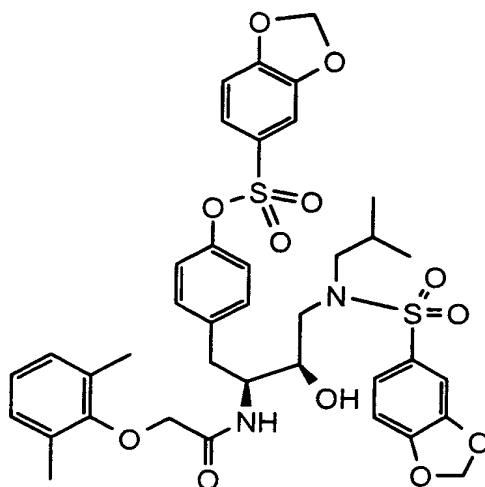
¹H-NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.20 (1H,m), 1.35 (1H,m), 1.91 (1H,m), 2.41 (1H,t), 2.60-2.80 (3H,m), 2.98 (2H,m), 3.20-3.40 (2H,m), 3.40-3.60 (4H,m), 3.66 (1H,t), 3.80 (1H,t), 4.80 (1H,dd), 5.04 (1H, s), 5.47 (1H,d), 6.13 (2H,s), 6.19 (2H,s), 6.90 (2H,d), 7.04 (2H,m), 7.10-7.40 (6H,m). MS: 777 (M+1)⁺.

15

20

Example 133

4-((2*S*,3*R*)-4-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-{[(2,6-dimethylphenoxy)acetyl]amino}-3-hydroxybutyl)phenyl 1,3-benzodioxole-5-sulfonate (350)

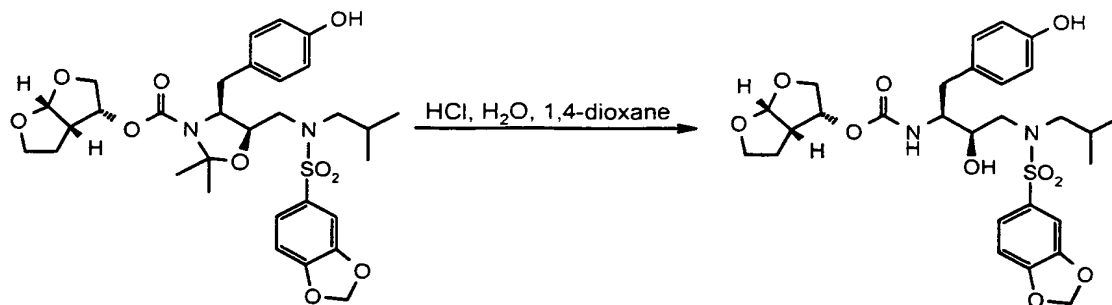


5

The coupling reaction was carried as described for the synthesis of *N*-{(1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(benzyloxy)benzyl]-2-hydroxypropyl}-2-(2,6-dimethylphenoxy)acetamide.

¹H-NMR (DMSO-*d*₆): δ 0.77 (3H,d), 0.81 (3H,d), 1.94 (1H,m), 2.07 (6H,s), 2.60-2.90 (3H,m), 2.90-3.10 (2H,m), 3.68 (1H,m), 3.81 (1H,d), 3.96 (1H,m), 4.05 (1H,d), 5.10 (1H,d), 6.12 (4H,m), 6.85-7.05 (7H,m), 7.15-7.35 (6H,m), 7.93 (1H,d). MS: 783 (M+1)⁺

Example 134



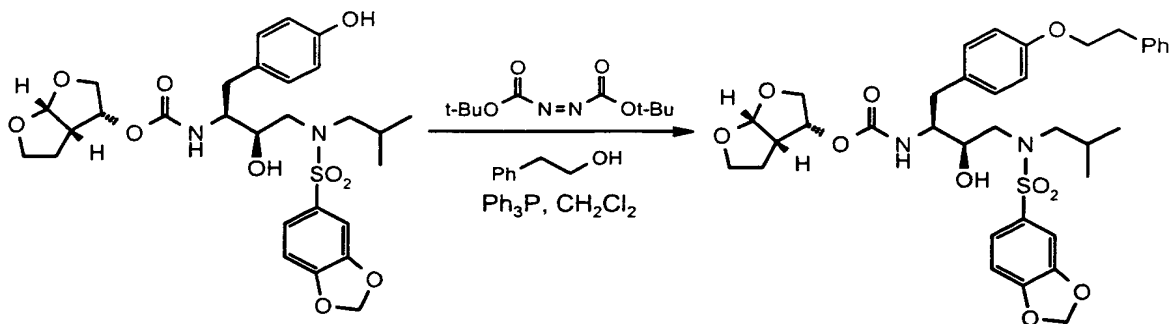
(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 5 hydroxy-1-(4-hydroxybenzyl)propylcarbamate (351)

A solution of 0.60 g (0.95 mmol) of

(3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 4-(4-hydroxybenzyl)-2,2-dimethyl-1,3-oxazolidine-3-

carboxylate in 5 mL of 1,4-dioxane was treated with 0.25
 mL of water followed by 5 mL of 4*N* HCl in 1,4-dioxane and
 the resulting solution was stirred at RT. After 1.5
 hours the solution was diluted with 25 mL of CH₂Cl₂ and
 the pH adjusted to approximately 12 by addition of 1*N*
 aqueous NaOH. The mixture was diluted with water and
 extracted with CH₂Cl₂ (3x). The combined CH₂Cl₂ extracts
 were washed with aqueous brine (3x), dried over MgSO₄, and
 concentrated *in vacuo*. The residue was subjected to
 flash chromatography (SiO₂, 8:2 EtOAc/hexane) to afford
 the desired compound as a white solid in 81% yield. ¹H NMR
 (CDCl₃): 7.29 (dd, 1H), 7.13 (s, 1H), 7.01 (d, 2H), 6.85
 (d, 2H), 6.70 (d, 2H), 6.05 (s, 2H), 5.61 (d, 1H), 5.01
 (m, 2H), 3.92 (dd, 1H), 3.80 (m, 4H), 3.68 (m, 2H), 3.08
 (dd, 1H), 2.91 (m, 5H), 2.81-2.62 (m, 2H), 1.80 (m, 1H),
 1.64 (m, 1H), 1.47 (m, 1H), 0.90 (d, 3H), 0.82 (d, 3H).
 MS (ESI): 593 (M+H).

Example 135



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 5 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-(phenethoxybenzyl)propylcarbamate (352)

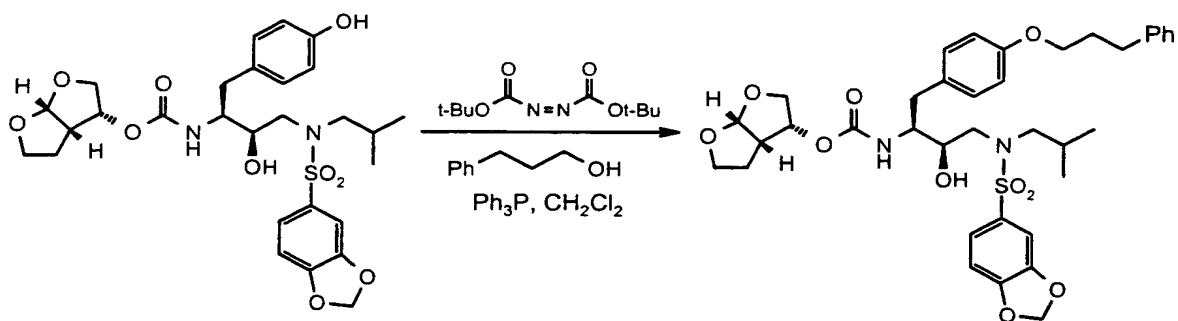
To a solution of 66 mg (0.25 mmol) of triphenylphosphine
 and 30 μ L (0.25 mmol) of phenethyl alcohol in 3 mL of
 anhydrous CH_2Cl_2 was added 58 mg (0.25 mmol) of di-tert-
 10 butyl azodicarboxylate. The resulting solution was
 stirred at RT for 5 minutes and was then treated with a
 solution of 50 mg (0.084 mmol) of

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-

15 hydroxy-1-(4-hydroxybenzyl)propylcarbamate in 2 mL of
 CH_2Cl_2 . After stirring at RT for 1.5 hours the solution
 was concentrated to dryness and the residue purified by
 flash chromatography (SiO_2 , 4:6 hexane/EtOAc) to give the
 desired product as a white foam in 72% yield. ^1H NMR

20 (CDCl_3): 7.34-7.17 (m, 6H), 7.15 (s, 1H), 7.07 (d, 2H),
 6.86 (d, 1H), 6.78 (d, 2H), 6.03 (s, 2H), 5.61 (d, 1H),
 4.98 (m, 1H), 4.86 (d, 1H), 4.09 (t, 2H), 3.92 (m, 1H),
 3.84-3.56 (m, 6H), 3.17-3.01 (m, 3H), 3.00-2.81 (m, 4H),
 2.80-2.64 (m, 2H), 1.78 (m, 1H), 1.56 (m, 1H), 1.47 (m,
 25 1H), 0.91 (d, 3H), 0.85 (d, 3H). MS(ESI): 697(M+H).

Example 136



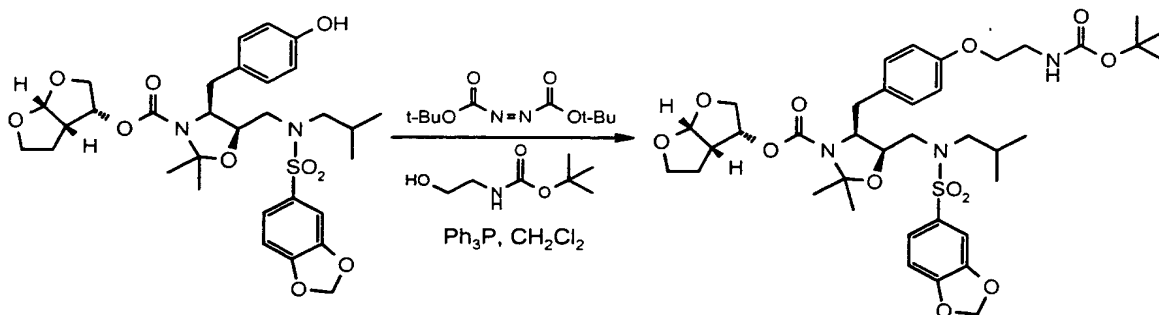
5 (3*R*,3*aS*,6*aR*)hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy
 -1-[4-([3-phenylpropyl]oxy)benzyl]propylcarbamate (353)

The title compound was prepared according to example 135
 with the exception that 3-phenyl-1-propanol was used

10 instead of phenethyl alcohol. ¹H NMR (CDCl₃): 7.33-7.11
 (m, 7H), 7.07 (d, 2H), 6.86 (d, 1H), 6.76 (d, 2H), 6.04
 (s, 2H), 5.61 (d, 1H), 4.99 (q, 1H), 4.86 (d, 1H), 3.97-
 3.72 (m, 7H), 3.65 (m, 2H), 3.09 (dd, 1H), 3.01-2.81 (m,
 4H), 2.80-2.64 (m, 4H), 2.06 (m, 2H), 1.85-1.40 (m, 3H),
 15 0.91 (d, 3H), 0.84 (d, 3H). MS(ESI): 711(M+H).

Example 137

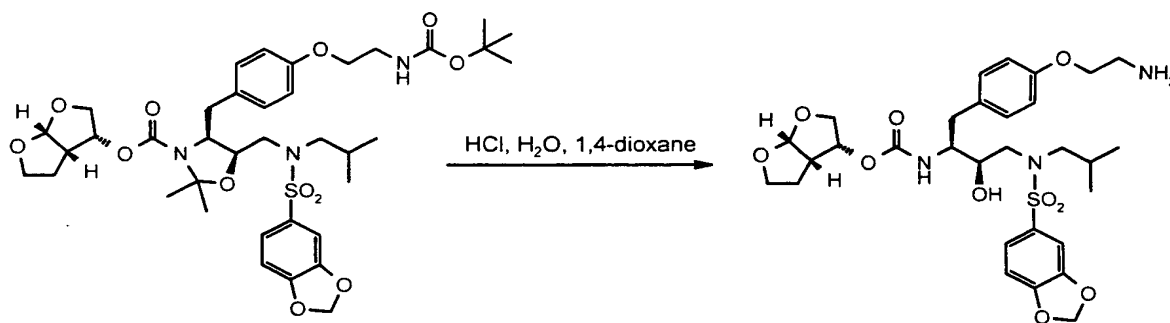
Step 1:



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (4S,5R)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 4-(4-[2-(tert-butoxycarbonylamino)ethoxy]benzyl)-2,2-
 dimethyl-1,3-oxazolidine-3-carboxylate

- 5 According to example 135 (with the exception that N-
 (tert-butoxycarbonyl)ethanolamine was used instead of
 phenethyl alcohol), (3R,3aS,6aR)hexahydrofuro[2,3-
 b]furan-3-yl (4S,5R)-5-{[(1,3-benzodioxol-5-
 ylsulfonyl)(isobutyl)amino]methyl}-4-(4-hydroxybenzyl)-
 10 2,2-dimethyl-1,3-oxazolidine-3-carboxylate was converted
 to the desired compound. MS(ESI): 798(M+Na).

Step 2:

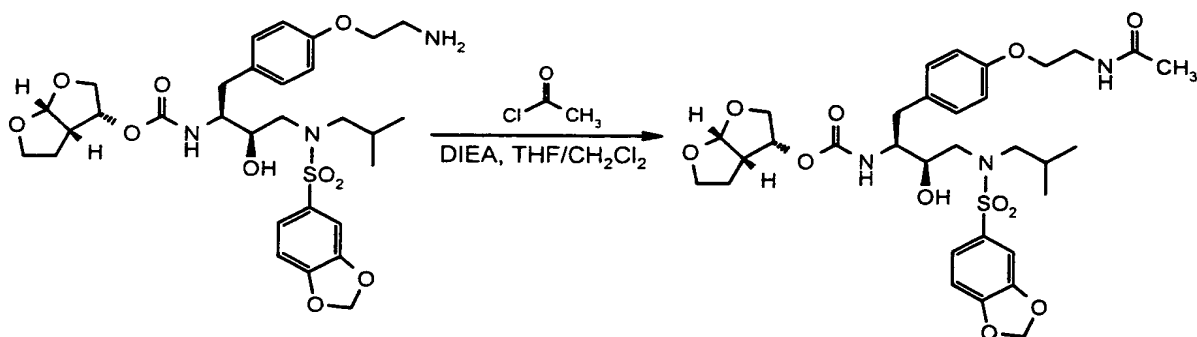


15

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-(2-aminoethoxy)benzyl]propylcarbamate (354)

- 20 The title compound was prepared according to example 134.
¹H NMR (CDCl₃): 7.39 (dd, 1H), 7.12 (s, 1H), 7.08 (d, 2H),
 6.84 (d, 1H), 6.77 (d, 2H), 6.03 (s, 2H), 5.61 (d, 1H),
 4.95 (m, 2H), 4.00-3.60 (m, 8H), 3.16-2.62 (m, 10H),
 2.20-1.40 (m, 5H), 0.90 (d, 3H), 0.82 (d, 3H). MS(ESI):
 25 636(M+H).

Example 138



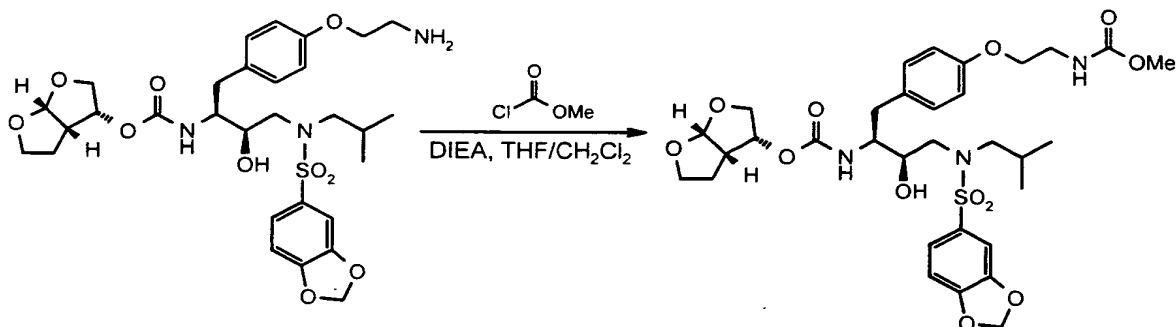
5 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-[4-(2-[acetylamino]ethoxy)benzyl]
propylcarbamate (355)

A solution of 21 mg (0.033 mmol) of

10 (3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
hydroxy-1-[4-(2-aminoethoxy)benzyl]propylcarbamate in 2
mL of 1:1 THF/CH₂Cl₂ was treated with 9 μL (0.050 mmol) of
N,N-diisopropylethylamine followed by 2.6 μL (0.036 mmol)
15 of acetyl chloride. The resulting solution was stirred
at RT. After 1.5 hours the solution was concentrated in
vacuo and the residue subjected to flash chromatography
(SiO₂, 95:5 CH₂Cl₂/MeOH) to afford the desired compound in
86 % yield as a white foam. ¹H NMR (CDCl₃): 7.29 (dd, 1H),
20 7.16-7.04 (m, 3H), 7.05 (d, 1H), 6.76 (d, 2H), 6.05 (s,
2H), 5.91 (br s, 1H), 5.01 (d, 1H), 5.05-4.89 (m, 2H),
3.94 (m, 3H), 3.80 (m, 3H), 3.72-3.51 (m, 5H), 3.08 (dd,
1H), 3.01-2.83 (m, 4H), 2.74 (m, 2H), 1.97 (s, 3H), 1.86-
1.45 (m, 3H), 0.90 (d, 3H), 0.84 (d, 3H). MS(ESI):

25 678 (M+H).

Example 139



5

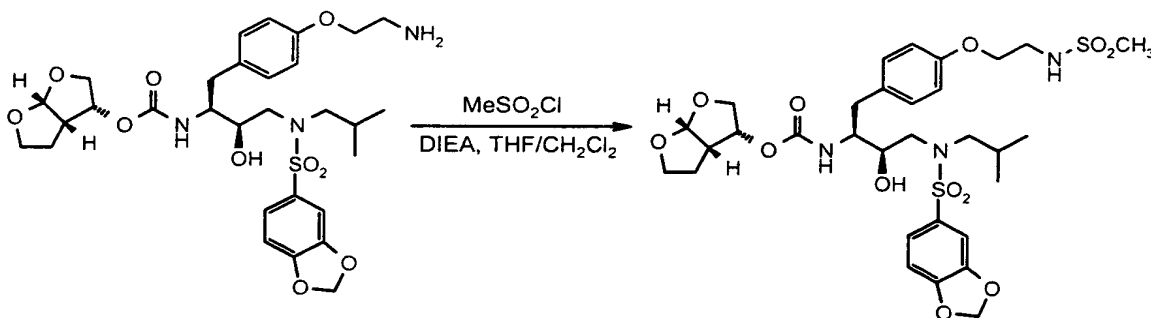
(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-(2-[(methoxycarbonyl)amino]ethoxy)

10 benzyl]propylcarbamate (356)

The title compound was prepared according to example 138
 with the exception that methyl chloroformate was used
 instead of acetyl chloride. ¹H NMR (CDCl₃): 7.29 (dd, 1H),
 7.16-7.02 (m, 3H), 6.84 (d, 1H), 6.76 (d, 2H), 6.04 (s,
 15 2H), 5.61 (d, 1H), 5.18-4.84 (m, 3H), 4.02-3.43 (m, 14H),
 3.09 (m, 1H), 2.91 (m, 4H), 2.72 (m, 2H), 1.85-1.43 (m,
 3H), 0.89 (d, 3H), 0.92 (d, 3H). MS(ESI): 694 (M+H).

Example 140

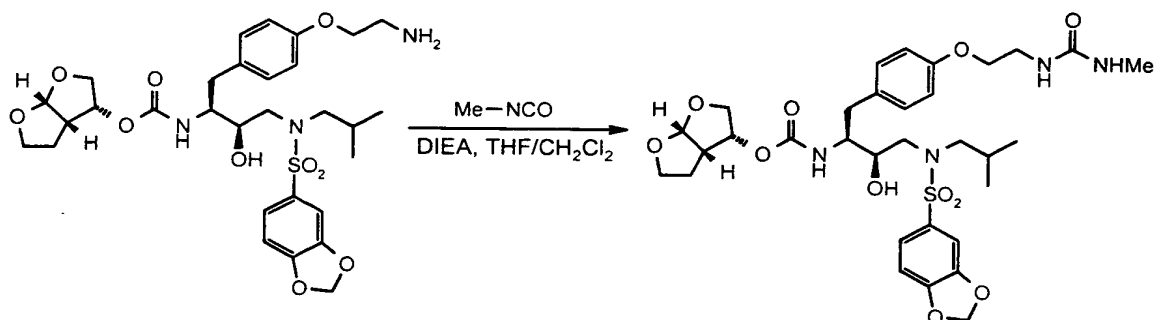
20



(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-(2-[(methylsulfonyl)amino]ethoxy)
 benzyl]propylcarbamate (357)

5 The title compound was prepared according to example 138
 with the exception that methanesulfonyl chloride was used
 instead of acetyl chloride. ¹H NMR (CDCl₃): 7.29 (dd, 1H),
 7.11 (m, 3H), 6.86 (d, 1H), 6.77 (d, 2H), 6.04 (s, 2H),
 5.61 (d, 1H), 4.98 (m, 2H), 4.84 (m, 1H), 4.09-3.44 (m,
 10 11H), 3.12-2.82 (m, 8H), 2.74 (m, 2H), 1.88-1.43 (m, 3H),
 0.89 (d, 3H), 0.81 (d, 3H). MS(ESI): 714 (M+H).

Example 141



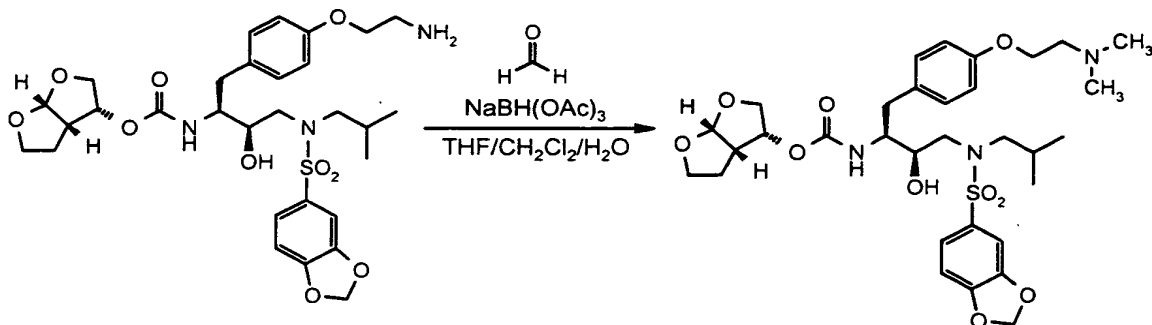
15

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-(2-[(methylamino]carbonyl)amino]
 20 ethoxy)benzyl]propylcarbamate (358)

The title compound was prepared according to example 138
 with the exception that methyl isocyanate was used
 instead of acetyl chloride.

¹H NMR (CDCl₃): 7.29 (dd, 1H), 7.13 (s, 1H), 7.08 (d, 2H),
 25 6.84 (d, 1H), 6.76 (d, 2H), 6.05 (s, 2H), 5.61 (d, 1H),
 5.10-4.90 (m, 2H), 4.04-3.48 (m, 11H), 3.15-2.67 (m,
 10H), 1.80 (m, 1H), 1.62 (m, 1H), 1.47 (m, 1H), 0.85 (m,
 6H). MS(ESI): 693 (M+H).

Example 142



5

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-(2-[dimethylamino]ethoxy)benzyl]
 propylcarbamate (359)

10

A solution of 21 mg (0.033 mmol) of

(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-(2-aminoethoxy)benzyl]propylcarbamate in 3

15

mL of 8:2 THF/CH₂Cl₂ was treated with 13 μL (0.17 mmol) of
 37% aqueous formaldehyde followed by 35 mg (0.17 mmol) of
 NaBH(OAc)₃ and the resulting cloudy solution was stirred
 at RT. After 3 hours the solution was filtered to remove
 solids and the filtrate concentrated *in vacuo*. The

20

residue was purified by flash chromatography (SiO₂, 95:5
 CH₂Cl₂/2M NH₃ in MeOH) to give the desired compound as a
 white foam in 68% yield. ¹H NMR (CDCl₃): 7.29 (dd, 1H),

7.13 (s, 1H), 7.07 (d, 2H), 6.85 (d, 1H), 6.80 (d, 2H),
 6.03 (s, 2H), 5.61 (d, 1H), 4.98 (q, 1H), 4.89 (d, 1H),

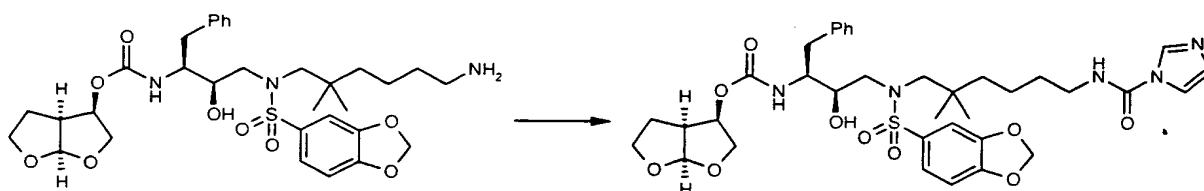
25

4.01 (t, 2H), 3.92 (m, 1H), 3.80 (m, 3H), 3.66 (m, 2H),
 3.09 (dd, 1H), 2.92 (m, 5H), 2.74 (m, 4H), 2.32 (s, 6H),

1.80 (m, 1H), 1.63 (m, 1H), 1.42 (m, 1H), 0.90 (d, 3H), 0.84 (d, 3H). MS(ESI): 664 (M+H).

Example 143

5 (3*R*,3*aS*,6*aR*)-Hexahydro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-[6-(*N'*-carbonylimidazol-1-yl)amino-2,2-dimethylhexyl][(3,4-methylenedioxyphenyl) sulfonyl]amino-2-hydroxypropyl)carbamate (360)



10 To a solution of (3*R*,3*aS*,6*aR*)-hexahydro[2,3-*b*]furan-3-yl *N*-((1*S*,2*R*)-1-benzyl-3-[6-amino-2,2-dimethylhexyl][(3,4-methylenedioxyphenyl) sulfonyl]amino-2-hydroxypropyl)carbamate (0.12 g, 0.19 mmol) in dichloromethane (1.5 mL) was added 1,1'-

15 carbonyldiimidazole (0.06 g, 0.37 mmol) and the mixture was stirred for 30 min at ambient temperature. The mixture was diluted with dichloromethane and washed with 5% citric acid/water (2X), saturated sodium bicarbonate/water, dried (sodium sulfate), evaporated,

20 and dried *in vacuo* to provide the title compound (0.14 g) as a solid foam; ¹H NMR (DMSO-*d*₆): 0.90 (6H, s), 1.12 (1H, dd), 1.20-1.35 (4H, m), 1.5 (2H, br s), 2.4 (1H, t), 2.7-2.8 (2H, m), 2.9-3.0 (2H, m), 3.2 (2H, dd), 3.25-3.35 (3H, m), 3.4 (1H, quartet), 3.5-3.6 (2H, m), 3.65 (1H, t), 3.75 (1H, quartet), 3.8 (1H, dd), 4.8 (1H, quartet),

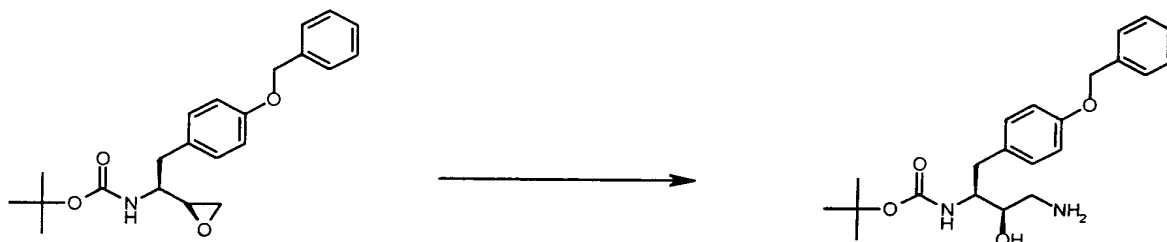
25 5.1 (1H, d), 5.5 (1H, d), 6.15 (2H, s), 6.98 (1H, s),

7.0 (1H, d), 7.10-7.25 (7H, m), 7.3 (1H, d), 7.6 (1H, s), 8.2 (1H, s), 8.4 (1H, t); MS: 742 (MH⁺); C₃₆H₄₇N₅O₁₀S.

Example 144

5 Step 1:

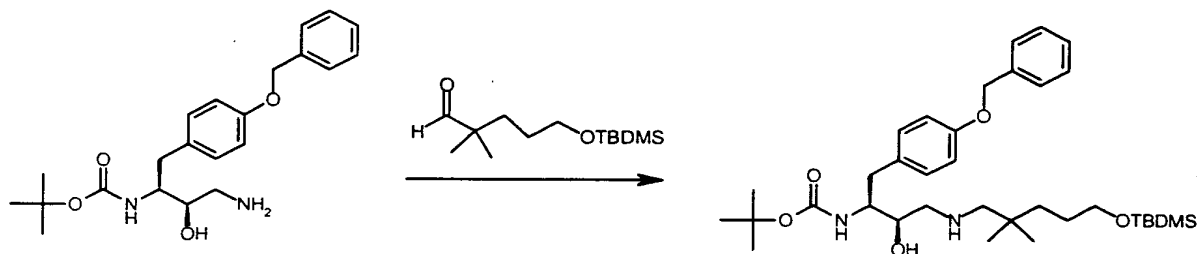
tert-Butyl N-(1*S*,2*R*)-1-benzyl-3-amino-1-[(4-benzyloxy)benzyl]-2-hydroxypropylcarbamate



- 10 To a saturated solution of ammonia in methanol (100mL) at 5 °C was added the epoxide (1.0g, 2.8mmol). Ammonia gas was continuously bubbled through the solution for an additional hour. The reaction was allowed to come to ambient temperature and stir for 48 hours. The methanol
- 15 was removed *in vacuo* and the resulting solid was triturated with ether (50mL), filtered and dried to afford the title compound (910mg, 85%) as a white solid.
- 20 ¹H NMR (DMSO-*d*₆): 1.22 (9H, s), 1.58 (2H, br), 2.41-2.53 (3H, m), 2.89 (1H, dd), 3.15-3.19 (1H, m), 3.33-3.45 (1H, m), 4.70 (1H, br), 5.00 (2H, s), 6.58 (1H, d), 6.83 (2H, d), 7.04 (2H, d), 7.25-7.39 (5H, m)

Step 2:

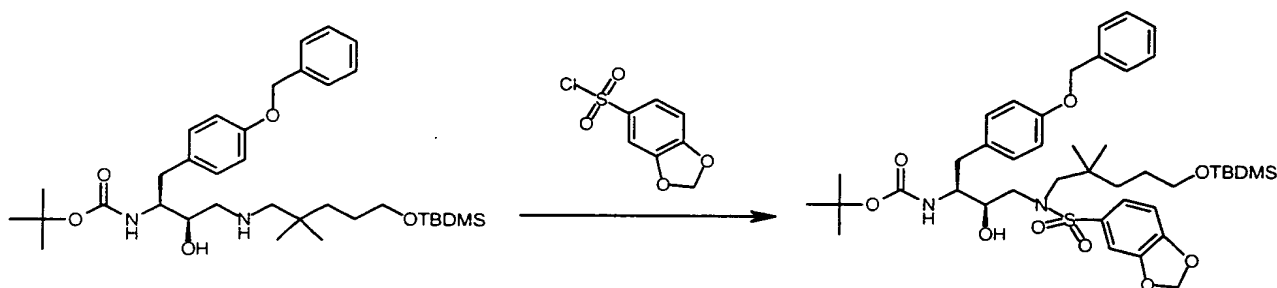
**tert-Butyl N-(1S,2R)-1-[(4-benzyloxy)benzyl]-3-[(5-tert-butyl-
dimethylsilyloxy-2,2-dimethyl)amino]-2-
hydroxypropylcarbamate**



- 5 To a stirred suspension of the above amine) (5.04g,
13.0mmol) in N,N-dimethylformamide (35mL), 1,2-
dichloroethane (35mL) and glacial acetic acid (.5mL) was
added a solution of 5-tert-butyl-2,2-dimethylpentanal (2.66g, 10.9mmol) in tetrahydrofuran
10 (35mL) over 15 minutes. Sodium triacetoxyborohydride
(2.76g, 13.0mmol) was added and the mixture was stirred
under nitrogen for 18 hours. The reaction was
concentrated *in vacuo* to approximately 1/3 volume, added
cold 2.5% sodium hydroxide (100mL) and extracted with
15 ethyl acetate (2x100mL). The combined ethyl acetate was
washed with water (2x50mL), dried (magnesium sulfate) and
concentrated. The residue was triturated with ether and
the resulting solid (amine starting material) was
filtered away. The filtrate was concentrated *in vacuo* to
20 afford the desired crude product (6.80g, 6.54 theory) as
a thick paste. The material was used without further
purification.

Step 3:

**tert-Butyl N-((1S,2R)-1-[(4-benzyloxy)benzyl]-3-(5-tert-butyl-
butyldimethylsilyloxy-2,2-dimethylpentyl)[(3,4-
methylenedioxyphenyl)sulfonyl]amino-2-
hydroxypropyl)carbamate**

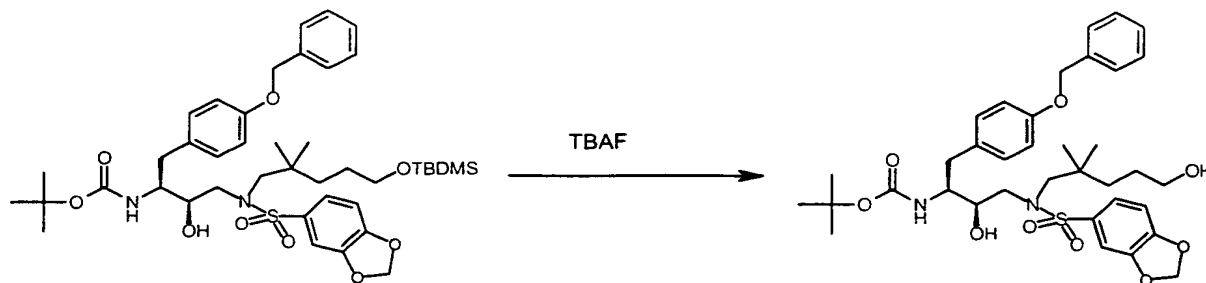


5

To a stirred solution of the crude amine from step 2, (6.80g), and N,N-diisopropylethylamine (3.7mL, 21.8mmol) in anhydrous dichloromethane (70mL) at 5 °C was added a solution of 3,4-methylenedioxyphenylsulfonyl chloride
10 (2.88g, 13.0mmol) in dichloromethane (30mL) over 15 minutes. The mixture was allowed to warm to ambient temperature and stir for 16 hours. The reaction was concentrated in vacuo, taken up in ethyl acetate (100mL), washed with water (50mL), 1N hydrochloric acid (2x50mL),
15 water (50mL), saturated sodium bicarbonate (2x50mL), dried (magnesium sulfate) and concentrated in vacuo to afford the desired crude product (8.55g) as a white foam. The material was used without further purification.

Step 4:

20 **tert-Butyl N-((1S,2R)-1-[(4-benzyloxy)benzyl]-3-(2,2-dimethyl-5-hydroxypentyl)[(3,4-methylenedioxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate**

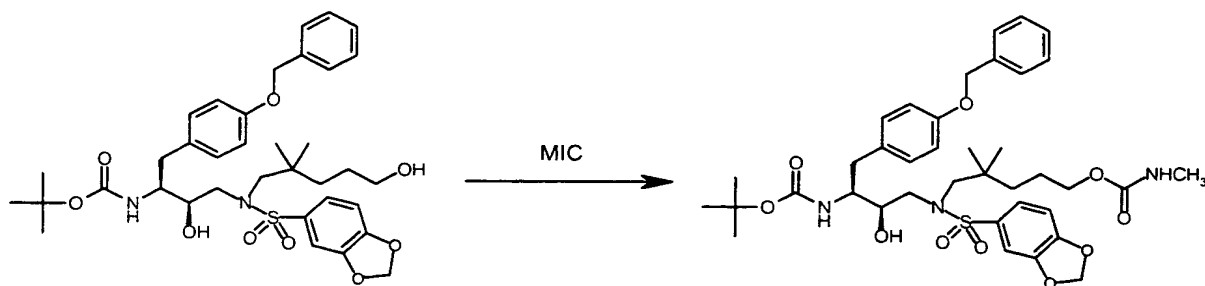


To a stirred solution of the crude product from step 3.) (8.5g, 10.6mmol) in tetrahydrofuran (85mL) was added tetrabutylammonium fluoride (13.8mL, 1M in tetrahydrofuran) over 15 minutes. After stirring for 18 hours, the reaction was concentrated *in vacuo*, added ether (100mL) and washed with water (3x50mL). The ether was dried (magnesium sulfate) and concentrated to a foam. Taken up in fresh ether (40mL) and stirred at ambient temperature for 120 hours. The resulting precipitate was filtered, rinsed with ether/hexane (1:1, 25mL) and dried to afford the title compound (3.2g, 44%) as a white solid. The filtrate was concentrated to a foam (2.95g). HPLC analysis showed approximately 65% product (not isolated). ¹H NMR (DMSO-d₆): 0.86 (6H, s), 1.11-1.23 (2H, m), 1.19 (9H, s), 1.30-1.35 (2H, m), 2.37 (1H, dd), 2.77-2.82 (2H, m), 2.91-2.97 (1H, m), 3.25-3.33 (5H, m), 3.61-3.67 (1H, m), 4.33 (1H, t), 4.89 (1H, d), 5.00 (2H, s), 6.11 (2H, s), 6.52 (1H, d), 6.82 (2H, d), 6.98-7.04 (3H, m), 7.22-7.38 (7H, m)

Step 5:

***tert*-Butyl N-((1*S*,2*R*)-1-[(4-benzyloxy)benzyl]-3-(2,2-dimethyl-5-*N'*-methylcarbamoyloxypentyl))[(3,4-**

methylenedioxyphenyl)sulfonyl] amino-2-hydroxypropyl) carbamate



To a stirred solution of the alcohol from step 4.)

5 (600mg, .88mmol) in dichloromethane (6mL) was added methylisocyanate (1.0mL, 17.50mmol). After stirring at ambient temperature for 48 hours, the reaction was concentrated *in vacuo* and purified by silica gel chromatography (1:1; ethyl acetate: hexane) to afford the

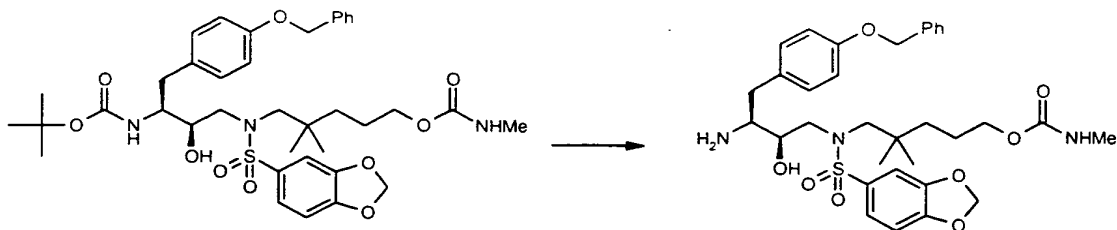
10 title compound (570mg, 88%) as a white foam. ¹H NMR (DMSO-d₆): 0.86 (6H, d), 1.18-1.23 (2H, m), 1.19 (9H, s), 1.43-1.49 (2H, m), 2.37 (1H, dd), 2.50 (3H, d), 2.77-2.82 (2H, m), 2.90-2.96 (1H, m), 3.28-3.31 (3H, m), 3.61-3.67 (1H, m), 3.84 (2H, t), 4.91 (1H, d), 5.00 (2H, s), 6.11 (1H, d), 6.52 (1H, d), 6.81-6.86 (3H, m), 6.98-7.04 (3H, m),

15 7.23-7.38 (7H, m)

Step 6:

*N*1-[(2*R*,3*S*)-3-amino-4-[(4-benzyloxy)phenyl]butyl]-*N*1-[2,2-dimethyl-5-(*N*2-methylcarbamoyloxy)pentyl]-3,4-methylenedioxy-1-benzenesulfonamide

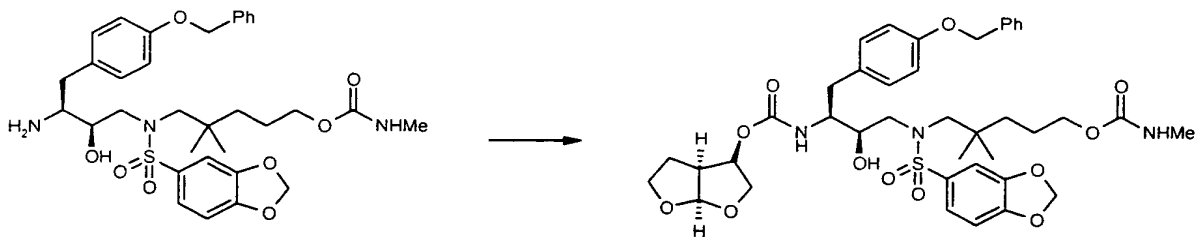
20



Treatment of the product from step 5 with trifluoroacetic acid/dichloromethane as previously described provided the title compound as a solid foam; ^1H NMR ($\text{DMSO}-d_6$): 0.8 (6H, d), 1.2-1.3 (2H, m), 1.4-1.5 (2H, m), 2.2 (1H, dd), 2.55 (3H, d), 2.6-2.7 (2H, m), 2.9 (1H, d), 3.1 (1H, dd), 3.2-3.3 (3H, m), 3.5 (2H, br d), 3.85 (2H, t), 4.6 (1H, d), 5.05 (2H, s), 6.1 (2H, s), 6.9 (3H, br d), 7.0 (1H, d), 7.08 (2H, br d), 7.2-7.4 (7H, m); MS: 642 (MH $^+$)

Step 7:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl N-((1S,2R)-1-[(4-benzyloxy)benzyl-3-[2,2-dimethyl-5-(N'-methylcarbamoyloxy)-pentyl]][(3,4-methylenedioxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate (361)



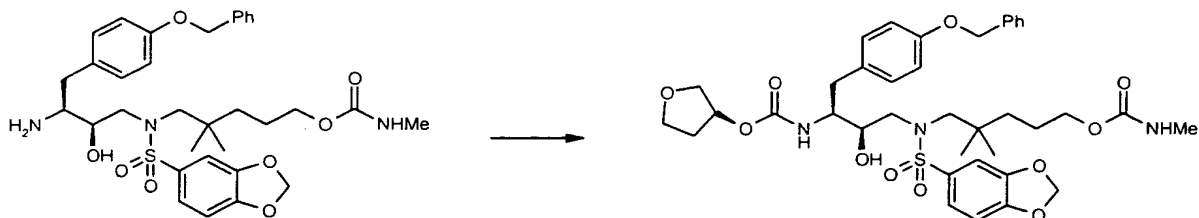
To a solution of the product from step 6 (0.11 g, 0.17 mmol) in acetonitrile (3 mL) was added

diisopropylethylamine (0.076 mL, 0.057 g, 0.44 mmol) and

[(3R,3aS,6aR)hexahydrofuro[2,3-b]furan-3-yl][4-nitrophenyl] carbonate (0.065 g, 0.22 mmol) and the mixture was stirred at ambient temperature for 24 h. Solvent was evaporated and the residue was dissolved in ethyl acetate, washed with saturated sodium bicarbonate/water (4X), dried (sodium sulfate), concentrated, and chromatographed (silica gel; hexanes/ethyl acetate) to provide the title compound as a solid foam; ¹H NMR (DMSO-d₆): 0.9 (6H, d), 1.1-1.3 (4H, m), 1.5 (2H, br s), 2.3 (1H, dd), 2.55 (3H, d), 2.7-2.8 (2H, m), 2.85 (2H, dd), 3.2-3.4 (4H, m), 3.5-3.6 (2H, m), 3.6-3.8 (2H, m), 3.82-3.90 (3H, m), 4.8 (1H, quartet), 5.0 (2H, s), 5.05 (1H, br d), 5.5 (1H, d), 6.1 (2H, s), 6.8-6.9 (3H, m), 7.0-7.1 (3H, m), 7.15-7.40 (7H, m); MS: 798 (MH⁺)

Example 145

(3S)-Tetrahydro-3-furanyl N-((1S,2R)-1-[(4-benzyloxy)benzyl-3-[2,2-dimethyl-5-(N'-methylcarbamoyloxy)-pentyl][(3,4-methylenedioxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate (362)

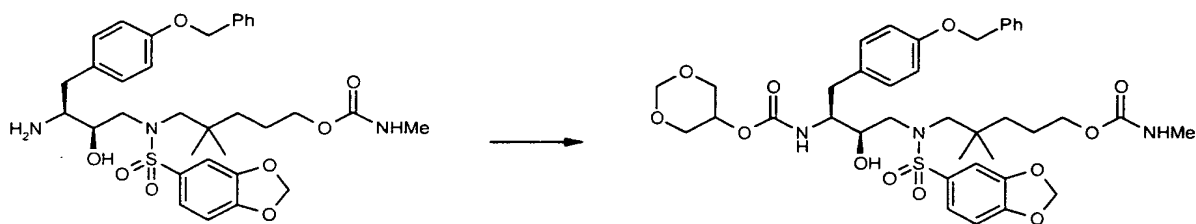


Treatment of the product from step 6 (Example 361) with (3S)-tetrahydro-3-furanyl N-succinimidyl carbonate as described in step g (example 801) provided the title compound as a solid foam; ¹H NMR (DMSO-d₆): 0.8 (6H, d),

1.05-1.10 (2H, m), 1.4-1.5 (2H, m), 1.7 (1H, br s), 1.95-2.05 (1H, m), 2.4 (1H, t), 2.55 (3H, d), 2.8 (2H, t), 2.9 (1H, dd), 3.2-3.35 (3H, m), 3.5-3.5 (4H, m), 3.8 (2H, br s), 4.9-5.0 (4H, m), 5.7 (1H, s), 6.2 (2H, s), 6.8-6.9 (3H, m), 7.00-7.15 (4H, m), 7.2-7.4 (7H, m); MS: 756 (MH+)

Example 146

1,3-Dioxan-5-yl *N*-((1*S*,2*R*)-1-[(4-benzyloxy)benzyl-3-[2,2-dimethyl-5-(*N'*-methylcarbamoyloxy)-pentyl]][(3,4-methylenedioxyphenyl)sulfonyl]amino-2-hydroxypropyl)carbamate (363)

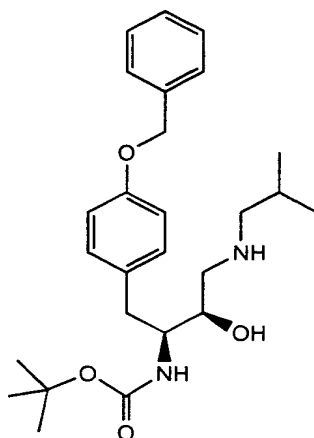


Treatment of the product from step 6 (example 361) with 1,3-dioxan-5-yl *p*-nitrophenyl carbonate as described in step 7 (example 627) provided the title compound as a solid foam; ¹H NMR (DMSO-*d*₆): 0.8 (6H, s), 1.1-1.3 (2H, m), 1.4-1.5 (2H, m), 2.4 (1H, t), 2.55 (3H, d), 2.75-2.85 (2H, m), 2.9 (1H, dd), 3.20-3.35 (2H, m), 3.45 (1H, d), 3.6-3.9 (6H, m), 4.3 (1H, br s), 4.65 (1H, br d), 4.75 (1H, br d), 4.9-5.0 (3H, m), 5.7 (1H, s), 6.1 (2H, s), 6.75-6.85 (3H, m), 6.95-7.05 (3H, m), 7.2-7.4 (8H, m); MS: 772 (MH+);

Example 147

Step 1:

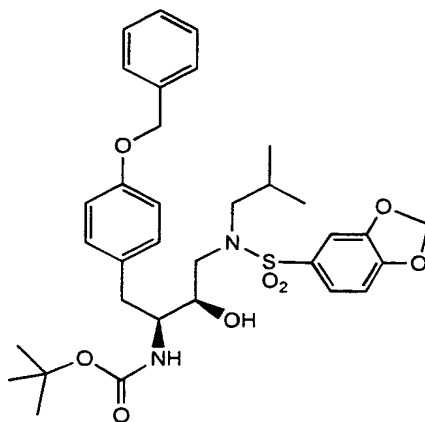
t-Butyl-(1S,2R)-1-(4-benzyloxy-benzyl)-3-i-butylamino-2-hydroxypropyl-carbamate



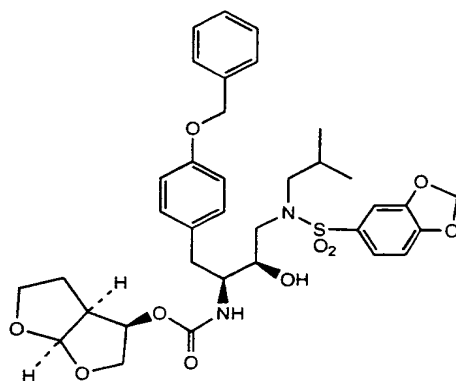
- 5 i-Butylamine (10.0 g, 137 mmol) was added to a solution of t-Butyl-(1S,2R)-1-(4-benzyloxy-benzyl)-2,3-epoxydo-propyl-carbamate (7.0 g, 18.9 mmol) (prepared according to the reference by Chen, P. at all., *Tetrahedron Letters*, Vol 38. p3175-8, 1997), in 100 ml i-propanol.
- 10 The mixture was stirred at 85°C for 2 hours, then it was cooled to 5°C and 500 ml water was added dropwise. The resulting suspension was stirred for 30 minutes at 5°C, then filtered. The solid was washed with water (3x100 ml) then dried under reduced pressure to obtain 8.3 g (99 %) title compound. ¹H -NMR: (CDCl₃): 0.88 (6H,d), 1.34 (9H, s), 1.68 (1H,m), 2.38 (2H,d), 2.64 (2H,d), 2.80 (1H,m), 2.86 (1H,dd), 3.40 (1H,m), 3.70 (2H, s(broad)), 4.63 (1H,d), 5.01 (2H,s), 6.88 (2H,d), 7.12 (2H,d), 7.40 (5H,m).

20 Step 2:

t-Butyl-N((1S,2R)-1-(4-benzyloxy-benzyl)-3-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carmamate



5 A solution of the product from Step 1 (8.3 g, 18.8 mmol), 3,4-methylenedioxysulfonylchloride (5.0 g, 22.7 mmol) and diisopropylethylamine (5.0 g, 38.7 mmol) stirred at 20°C for 4 hours. 200 ml water was then added
10 to the reaction mixture and the phases were separated. The aqueous phase was extracted with dichloromethane (3x100 ml), then the organic phases were combined, dried with magnesium sulfate, filtered and concentrated. The residue was dissolved in 300 ml ether, 50 g silicagel was
15 added to the solution, then the mixture was filtered. The filtrate was concentrated under reduced pressure, and the residue was mixed with 300 ml hexane. The mix was stirred for three hours at 20°C, then the solid was filtered and dried to afford 10.9 g (93 %) title compound. ¹H NMR
20 (CDCl₃): 0.85 (3H,d), 0.88 (3H,d), 1.34 (9H,s), 1.82 (1H,m), 3.04 (2H,m), 3.68 (1H,s(broad)), 3.75 (1H,s(broad)), 3.86 (1H,s), 4.60 (1H,d), 5.02 (2H,s), 6.04 (2H,s), 6.88 (3H,m), 7.15 (3H,m), 7.35 (6H,m).



Step 3:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(4-benzyloxy-benzyl)-3-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carbamate (364)

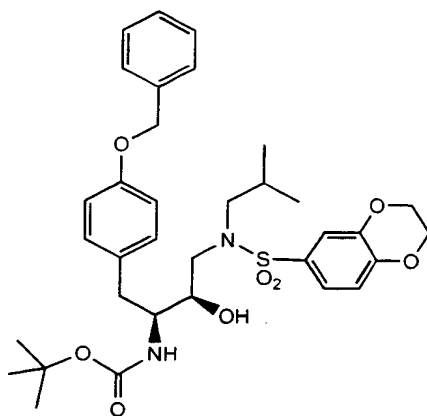
100 ml trifluoroacetic acid was added dropwise to a solution of the product from Step 2 (10.2g, 16.3 mmol) in 200 ml dichloromethane. The mixture was stirred for 1 hour at 20°C, then the solvents were evaporated under reduced pressure. The residue was dissolved in 200 ml dichloromethane and 300 ml 5% aqueous sodium carbonate solution was added, then the mixture was stirred at 20°C for 15 minutes. Phases were separated, then the aqueous phase was extracted with additional (2x100 ml) dichloromethane. Organic phases were combined then concentrated under reduced pressure. The residue was dissolved in 150 ml acetonitrile. Diisopropylethylamine (8.0 g, 62 mmol) and (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-4-nitrophenyl carbonate (8.0 g, 27.1 mmol) was added, and the solution was stirred for 12 hours at 20°C. 10 ml 25% aqueous ammonium hydroxide solution was added, then the mixture was stirred for an additional hour. The solvents were removed under reduced pressure,

the residue dissolved in 500 ml ether, and the solution was extracted with 5% sodium carbonate solution (10x100 ml). The organic phase was dried with anhydrous sodium carbonate, filtered and concentrated under reduced pressure. 20 ml ether was added to the residue, then after a small amount of solid formation 200 ml hexane was added, and the mixture was stirred at 20°C for 1 hour, then filtered and dried, to obtain 11.3 g (quant.) title compound. ¹H NMR (DMSO-d₆): 0.76 (3H,d), 0.82 (3H,d), 1.2 (1H,m), 1.35 (1H,m), 1.92 (1H,m), 2.35 (1H,t), 2.70 (3H,m), 2.85-3.05 (2H,m), 3.45 (1H,m), 3.55 (3H,m), 3.66 (1H,t), 3.80 (1H,dd), 4.81 (1H,m), 5.00 (3H,s(broad)), 5.47 (1H,d), 6.13 (2H,s), 6.82 (2H,d), 7.06 (3H,m), 7.20-7.40 (7H,m); MS: 682 (M+).

Example 148

Step 1:

t-Butyl-N((1*S*,2*R*)-1-(4-benzyloxy-benzyl)-3-*i*-butyl-[(3,4-ethylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carmamate

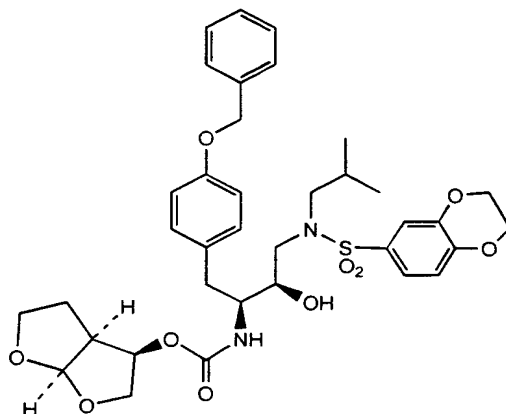


The reaction was carried out as described in Step 2, Example 147, except 3,4-ethylenedioxyphenyl sulfonylchloride was used in the reaction (83%).

¹H NMR (CDCl₃): 0.86 (3H,d), 0.89 (3H,d), 1.34 (9H,s),
 5 1.82 (1H,m), 2.85 (4H,m), 3.04 (2H,m), 3.69
 (1H,s(broad)), 3.76 (1H,s(broad)), 3.91 (1H,s), 4.27 (4H,
 m), 4.61 (1H,d), 5.03 (2H,s), 6.89 (2H,d), 6.93 91H,d),
 7.14 (2H,d), 7.20-7.45 (7H, m).

Step 2:

10 (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-(
 (4-benzyloxy-benzyl)-3-i-butyl-[(3,4-
 ethylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl-
 caramate (365)



15 The reaction was carried out as described in Step3,
 Example 147 except the product from Step 1, Example 144
 was used in the reaction (43%). ¹H NMR (DMSO-d₆): 0.78
 (3H,d), 0.84 (3H,d), 1.20 (1H,m), 1.35 (1H,m), 1.95
 (1H,m), 2.37 (1H,t), 2.70 (3H,m), 2.95 (2H,m), 3.45
 20 (1H,m), 3.58 (3H,m), 3.68 (1H,t), 3.82 (1H,m), 4.28
 (4H,d), 4.82 (1H,m), 5.01 (3H, s+m), 5.48 (1H,d), 6.84

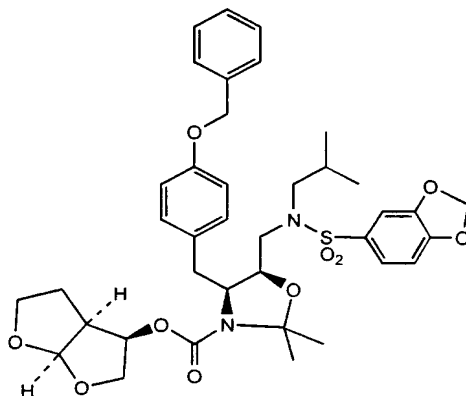
000090" 4946560

(2H,d), 7.02 (1H,d), 7.08 (2H, d), 7.20-7.40 (7H,m); MS: 696 (M+).

Example 149

5 Step 1:

N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-, (4S,5R)-4-(4-benzyloxy-benzyl)-5-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-aminomethyl-2,2-dimethyl-oxazolidine



10

2,2-dimethoxy-propane (20 g, 192 mmol) and p-toluenesulfonic acid (1.0 g, 5.8 mmol) was added to the solution of the product of Example 630 (11.3 g, 16.5 mmol) in 200 ml dichloromethane. The solution was
15 refluxed for 4 hours, then cooled to 20°C. The mixture was extracted with 5% sodium carbonate solution, then the organic phase was dried with magnesium sulfate, filtered and concentrated under reduced pressure. The residue was
20 purified on silicagel using hexane-ethyl acetate (1:1) as eluent, to provide 7.78 g (60 %) title compound. ¹H NMR (CDCl₃): 0.83 (3H,d), 0.91 (3H,d), 1.40, 1.48 (3H,s)*, 1.56, 1.64 (3H,s)*, 1.85 (2H,m), 2.0 (1H,m), 2.70 (3H,m),

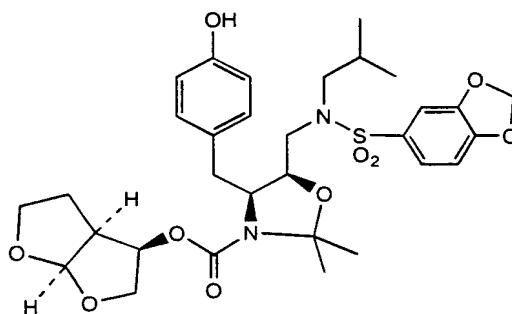
2.80 (1H,m), 3.00 (3H,m), 3.40 (2H,m), 3.80 (2H,m), 3.95 (1H,m), 4.20 (1H,m), 4.30 (1H,m), 4.89 (1H,dd), 5.03 (2H,s), 5.65, 5.68 (1H,d)*, 6.00 (2H,s), 6.79 (1H,d), 6.89 (2H,d), 7.02 (2H,d), 7.10 (1H,m), 7.30-7.45 (6H,m).

5 *: possible indication for rotamers.

Step 2:

N- (3R,3aS,6aR) -Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-, (4S,5R)-4-(4-hydroxybenzyl)-5-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-aminomethyl-2,2-dimethyl-oxazolidine (366)

10



8 g palladium (on charcoal, 10% Pd, Degussa type) was added to a stirred solution of the product of Step 1 (7.7 g, 10.6 mmol) in 400 ml tetrahydrofuran. The mixture was stirred under atmospheric pressure of hydrogen for 12 hours. The catalyst was filtered, and the solvent was removed under reduced pressure.

15

The residue was stirred for 2 hours in 200 ml hexane, then the solid was filtered, washed with hexane (2x20 ml) to obtain 6.4 g (95 %) title compound. ¹H NMR (CDCl₃): 0.83 (3H,d), 0.91 (3H,d), 1.41, 1.48 (3H,s)*, 1.56, 1.65 (3H,s)*, 1.85 (2H,m), 2.00 (1H,m), 2.60-3.10 (6H,m), 3.40 (2H,m), 3.70-4.35 (6H,m), 4.90 (1H, dd), 5.05-5.30

20

(2H,m), 5.66, 5.69 (1H,d)*, 6.06 (2H,s), 6.75 (2H,d),
6.83 (1H,d), 6.97 (2H,d), 7.00-7.15 (2H,m); MS: 632 (M+).
*: possible indication for rotamers.

5

Example 150

Step 1:

General procedure for the aralkylation of N-(3R,3aS,6aR)-
Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-, (4S,5R)-4-
(4-hydroxybenzyl)-5-i-butyl-[(3,4-methylenedioxyphenyl)
10 sulfonyl]-aminomethyl-2,2-Dimethyl-oxazolidine

0.5 mmol aralkyl halide[1] was added to a stirred mixture
of N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-
oxycarbonyl-, (4S,5R)-4-(4-hydroxybenzyl)-5-i-butyl-
[(3,4-methylenedioxyphenyl)sulfonyl]-aminomethyl-2,2-
15 Dimethyl-oxazolidine (126 mg, 0.2 mmol) and cesium
carbonate (250 mg, 0.76 mmol) in 2 ml N,N-
dimethylformamide. The mixture was stirred at the
indicated temperature[2] for indicated number of hours[3]
then diluted with 100 ml ether at 20°C. The mixture was
20 filtered, and the filtrate was extracted with water
(10x20 ml). The organic phase was dried with magnesium
sulfate, filtered and concentrated under reduced pressure
to obtain the following compounds:

N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-
25 oxycarbonyl-, (4S,5R)-4-{4-[3-(4-fluorophenyl)-1-

propyloxy]-benzyl}-5-i-butyl-[(3,4-methylenedioxyphenyl)
sulfonyl]-aminomethyl-2,2-dimethyl-oxazolidine

5

[1]: 1-chloro-3-(4-fluorophenyl)-propane; [2]: 60°C; [3]:
6 hours; MS: 768 (M+).

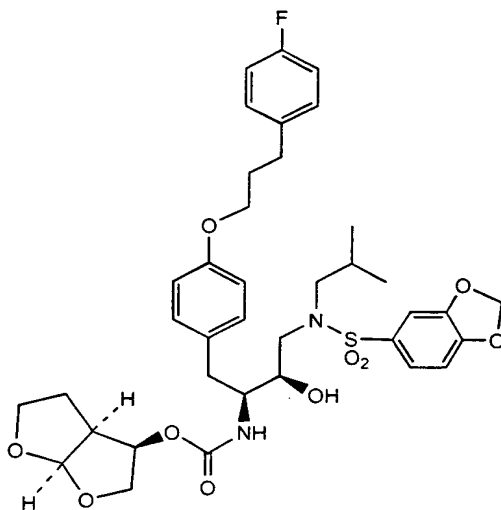
Step 2:

General procedure for the deprotection of N-(3R,3aS,6aR)-
10 Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-, (4S,5R)-4-
(4-aralkyloxybenzyl)-5-i-butyl-[(3,4-
methylenedioxyphenyl)sulfonyl]-aminomethyl-2,2-Dimethyl-
oxazolidine (the product of Step 1.)

4 M HCl in dioxane (7.5 ml) and water (0.2 g) was added
15 to a stirred solution of the product of Step 1 in 15 ml
dioxane. The solution was stirred for 5 hours, then the
solvents were removed under reduced pressure. The residue
was dissolved in 100 ml ethyl acetate, washed with 5%
aqueous sodium bicarbonate solution, then the organic
20 phase dried with magnesium sulfate. The solvents were
removed and the residue was purified either with
filtration from silicagel slurry and crystallization(a)

or silicagel chromatography(b), using the eluent(s) as indicated [1] to obtain the following compounds (yield:[2]):

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-
5 {4-[3-(4-fluorophenyl)-1-propyloxy]-benzyl}-3-i-butyl-
[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-
hydroxypropyl-carbamate (366)



[1]: b, Hexane-EtOAc (1:1) then Hexane-EtOAc (1:2); [2] :
10 36 %. ¹H-NMR (DMSO-*d*₆): 0.76 (3H,d), 0.81 (3H,d), 1.15
(1H,m), 1.31 (1H,m), 1.93 (3H,m), 2.34 (1H,t), 2.68
(5H,m), 2.85-3.00 (3H,m), 3.40-3.90 (8H,m), 4.81 (1H,dd),
4.98 (1H,d), 5.47 (1H,d), 6.13 (2H,s), 6.74 (2H,d), 7.04
(4H,m), 7.20 (4H,m), 7.27 (1H,d); MS: 728 (M+).

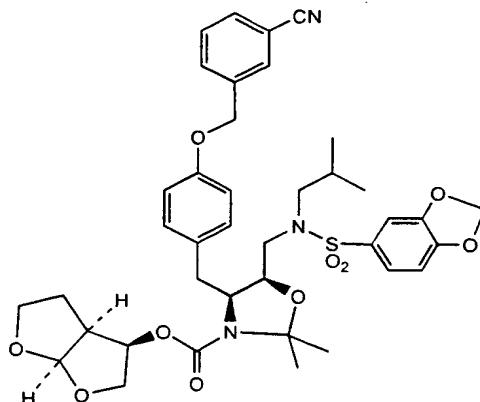
15

Example 151

Step 1:

N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-
oxycarbonyl-, (4S,5R)-4-[4-(3-cyanobenzyloxy)-benzyl]-5-

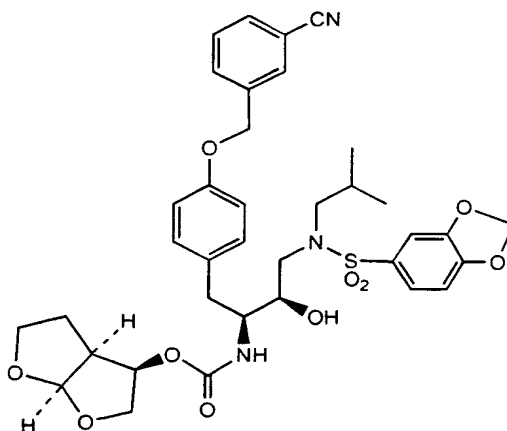
i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-aminomethyl-
2,2-dimethyl-oxazolidine



5 [1]: 3-cyanobenzyl bromide; [2]: 20°C; [3]: 1 hour; MS:
747 (M+).

Step 2:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-
10 [4-(3-cyanobenzyloxy)-benzyl]-3-i-butyl-[(3,4-
methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl-
carbamate (367)



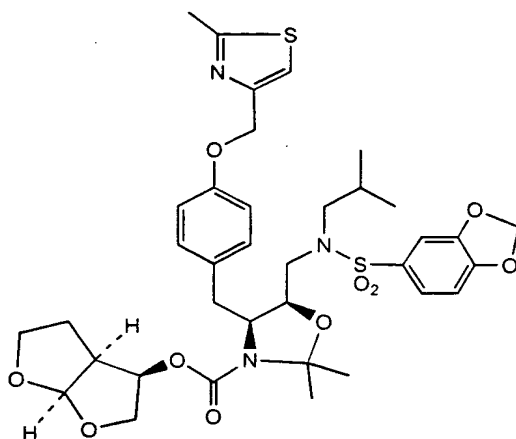
[1]: b, Hexane-EtOAc (1:1) then Hexane-EtOAc (1:2); [2]:
 46 %. ¹H NMR (DMSO-d₆): 0.75 (3H,d), 0.81 (3H,d), 1.19
 (1H,m), 1.31 (1H,m), 1.92 (1H,m), 2.35 (1H,t), 2.41
 (3H,m), 2.85-3.00 (3H,m), 3.40-3.60 (5H,m), 3.65 (1H,t),
 5 3.80 (1H,dd), 4.81 (1H,dd), 4.99 (1H,d), 5.07 (2H,s),
 5.47 (1H,d), 6.13 (2H,s), 6.83 (2H,d), 7.04 (3H,m), 7.20
 (2H,m), 7.27 (1H,d), 7.56 (1H,t), 7.75 (2H,t), 7.85 (1H,
 s); MS: 707(M+).

10

Example 152

Step 1:

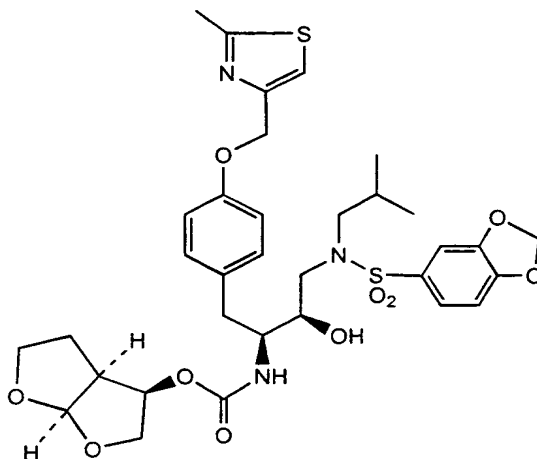
N- (3R,3aS,6aR) -Hexahydrofuro[2,3-b]furan-3-yl-
 oxycarbonyl-, (4S,5R) -4- [4- (2-methylthiazolo-4-
 methyloxy) -benzyl] -5-i-butyl- [(3,4-
 15 methylenedioxyphenyl) sulfonyl] -aminomethyl-2,2-dimethyl-
 oxazolidine



[1]: 4-chloromethyl-2-methylthiazole hydrochloride; [2]:
 20 70°C; [3]: 5 hour; MS: 743 (M+).

Step 2:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-[4-(2-methylthiazolo-4-methyloxy)-benzyl]-3-i-butyl-
5 [(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carbamate (368)



[1]: b, EtOAc (neat); [2]: 36 %. ¹H NMR (DMSO-d₆): 0.76 (3H,d), 0.82 (3H,d), 1.20 (1H,m), 1.33 (1H,m), 1.92 (1H,m), 2.35 (1H,t), 2.61 (3H,s), 2.65 (3H,m), 2.85-3.00 (3H,m), 3.40-3.60 (5H,m), 3.68 (1H,t), 3.80 (1H,dd), 4.81 (1H,dd), 4.99 (3H,s(broad)), 5.47 (1H,d), 6.13 (2H,s), 6.83 (2H,d), 7.06 (3H,m), 7.20 (2H,m), 7.27 (1H,d), 7.47 (1H,s); MS: 703 (M⁺).

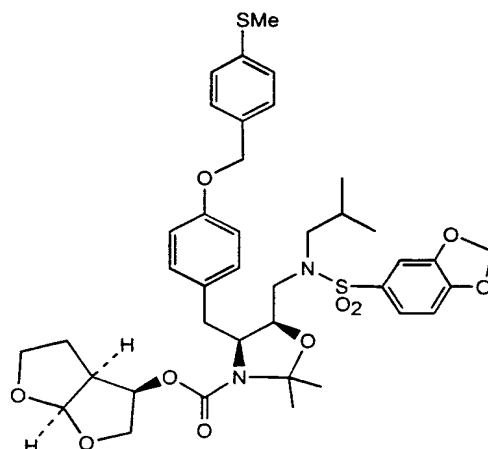
15

Example 153

Step 1:

N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-, (4S,5R)-4-[4-(4-methylthio-benzoyloxy)-

benzyl]-5-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-
aminomethyl-2,2-dimethyl-oxazolidine

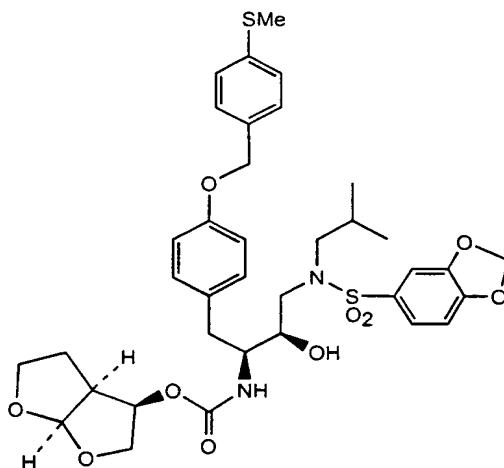


5

[1]: 4-methylthiobenzylchloride; [2]: 40°C; [3]: 4 hour;
MS: 768 (M+).

Step 2:

10 (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-
[4-(4-methylthio-benzyloxy)-benzyl]-3-i-butyl-[(3,4-
methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl-
carmamate (369)



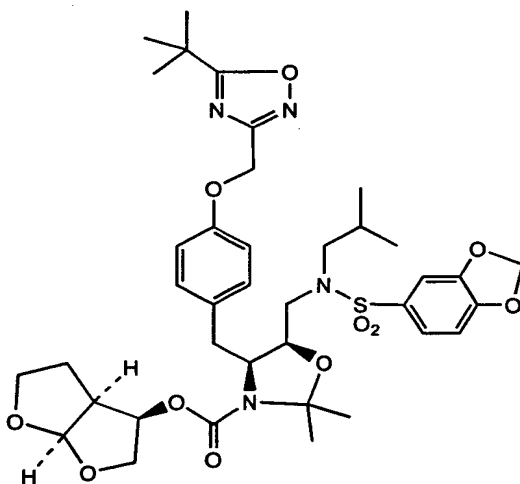
[1]: b, Hexane-EtOAc (1:1) then Hexane-EtOAc (1:2); [2]:
 17 %. ¹H NMR (DMSO-d₆): 0.75 (3H,d), 0.81 (3H,d), 1.20
 (1H,m), 1.30 (1H,m), 1.92 (1H,m), 2.35 (1H,t), 2.42
 5 (3H,s), 2.70 (3H,m), 2.85-3.00 (3H,m), 3.45 (1H,m), 3.55
 (3H,m), 3.66 (1H,t), 3.80 (1H,dd), 4.81 (1H,dd), 4.98
 (3H,s+m), 5.47 (1H,d), 6.12 (2H,s), 6.80 (2H,d), 7.05
 (3H,m), 7.15-7.35 (6H,m); MS: 728 (M+).

10

Example 154

Step 1:

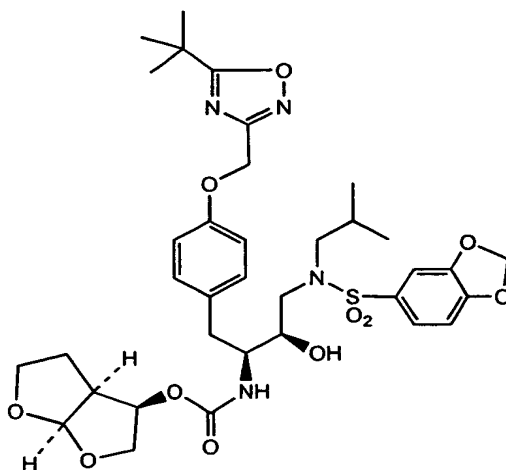
N- (3R,3aS,6aR) -Hexahydrofuro [2,3-b] furan-3-yl-
 oxycarbonyl-, (4S,5R) -4- [4- (5-t-butyl-1,2,4-oxadiazolo-3-
 methyloxy) -benzyl] -5-i-butyl- [(3,4-
 15 methylenedioxyphenyl) sulfonyl] -aminomethyl-2,2-dimethyl-
 oxazolidine



[1]: 5-t-butyl-3-chloromethyl-1,2,4-oxadiazole; [2]:
 20°C; [3]: 12 hour; MS: 770 (M+).

Step 2:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-[4-(5-t-butyl-1,2,4-oxadiazolo-3-methyloxy)-benzyl]-3-i-
5 butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl-carbamate (370)



[1]: b, Hexane-EtOAc (1:1) then Hexane-EtOAc (1:2); [2] :
10 56 %. ¹H NMR (DMSO-d₆): 0.76 (3H,d), 0.83 (3H,d), 1.19
(1H,m), 1.30 (1H,m), 1.36 (9H,s) 1.93 (1H,m), 2.37
(1H,t), 2.72 (3H,m), 2.90-3.02 (3H,m), 3.42-3.60 (4H,m),
3.68 (1H,t), 3.82 (1H,dd), 4.81 (1H,dd), 5.0
(1H,s(broad)), 5.12 (2H,s), 5.47 (1H,d), 6.13 (2H,s),
15 6.85 (2H,d), 7.03 (1H,d), 7.10 (2H,d), 7.20 (2H,d), 7.28
(1H,d); MS: 730 (M+).

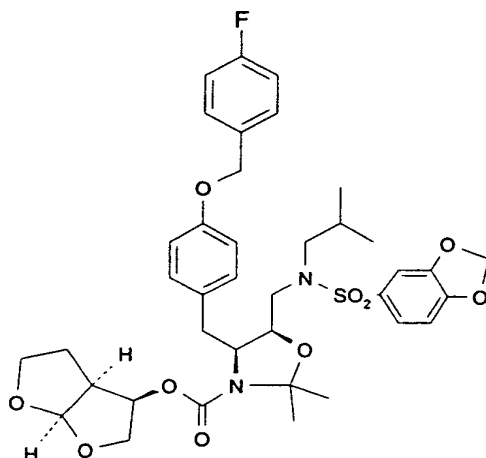
Example 155

Step 1:

N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-
20 oxycarbonyl-, (4S,5R)-4-[4-(4-flouorobenzyloxy)-benzyl]-5-

i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-aminomethyl-
2,2-dimethyl-oxazolidine (371)

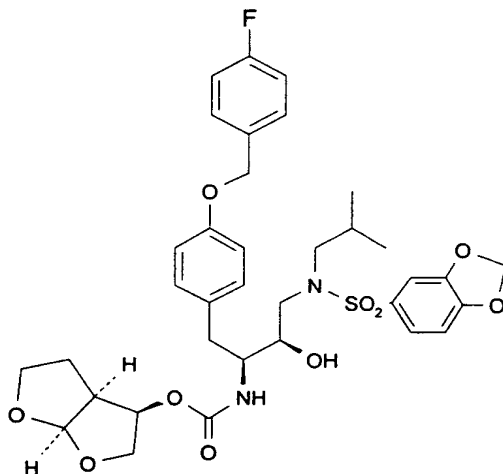
[1]: 4-fluorobenzylbromide; [2]: 20°C; [3]: 1 hour; MS:



741 (M+).

5 Step 2:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-[4-(4-fluorobenzoyloxy)-benzyl]-3-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carbamate (371)



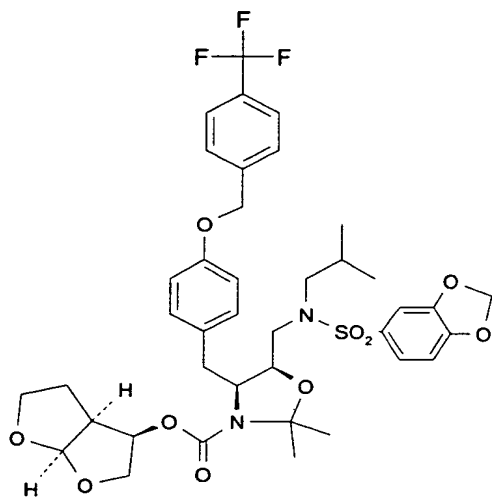
10 [1]: a, Hexane-EtOAc (1:1); [2]: 41%. ¹H-NMR (DMSO-d₆): 0.75 (3H,d), 0.81 (3H,d), 1.18 (1H,m), 1.31 (1H,m), 1.90 (1H,m), 2.34 (1H,t), 2.62-2.74 (3H,m), 2.89-3.02 (2H,m),

3.40-3.49 (1H,m), 3.50-3.60 (3H,m), 3.65 (1H,m), 3.80
(1H,dd), 4.80 (1H,dd), 4.97 (2H,s(broad)), 4.99 (1H,d),
5.46 (1H,d), 6.12 (2H,s), 6.81 (2H,d), 7.02 (1H,d), 7.06
(2H,d), 7.15 (1H,d), 7.18-7.23 (3H,m), 7.27 (1H,dd), 7.43
5 (2H,dd); MS: 701 (M+).

Example 156

Step 1:

N- (3R,3aS,6aR) -Hexahydrofuro [2,3-b] furan-3-yl-
oxycarbonyl-, (4S,5R) -4- [4- (4-trifluoromethylbenzyloxy) -
10 benzyl] -5-i-butyl- [(3,4-methylenedioxyphenyl) sulfonyl] -
aminomethyl-2,2-dimethyl-oxazolidine

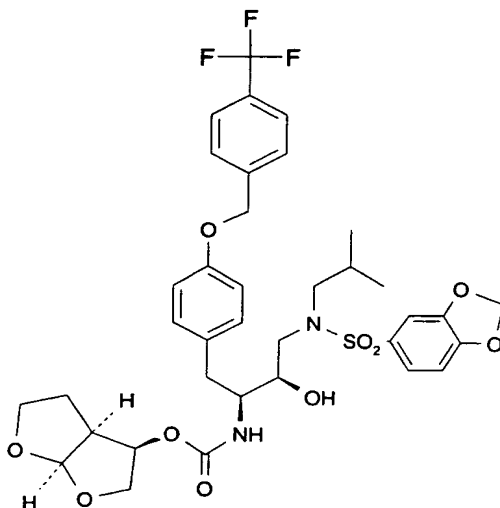


[1]: 4-trifluoromethylbenzylbromide; [2]: 20°C; [3]: 1
hour; MS: 791 (M+).

15

Step 2:

(3R,3aS,6aR) -Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-[4-(4-trifluoromethylbenzyloxy)-benzyl]-3-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carmamate (372)



5

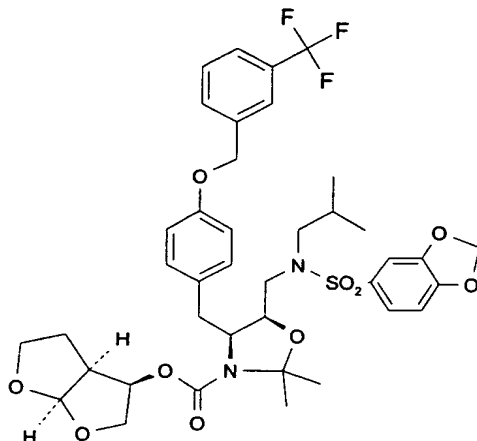
[1]: a, Hexane-EtOAc (1:1); [2] : 47 %. ¹H-NMR (DMSO-*d*₆): 0.75 (3H,d), 0.81 (3H,d), 1.16 (1H,m), 1.26 (1H,m), 1.91 (1H,m), 2.34 (1H,t), 2.65-2.76 (3H,m), 2.89-3.00 (2H,m), 3.45 (1H,m), 3.54 (3H,m), 3.63 (1H,m), 3.79 (1H,dd), 4.80 (1H,dd), 4.99 (1H,d), 5.12 (2H,s(broad)), 5.44 (1H,d), 6.12 (2H,s), 6.83 (2H,d), 7.02 (1H,d), 7.07 (2H,d), 7.21 (2H,d), 7.27(1H,d), 7.60 (2H,d), 7.70 (2H,d); MS: 751 (M+).

10

Example 157

15 Step 1:

N-(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-, (4S,5R)-4-[4-(3-trifluoromethylbenzyloxy)-benzyl]-5-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-aminomethyl-2,2-dimethyl-oxazolidine

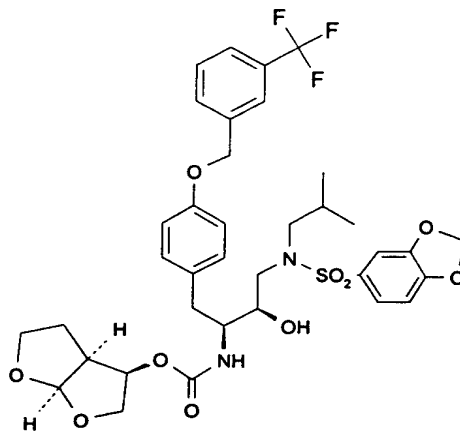


5

[1]: 3-trifluoromethylbenzylbromide; [2]: 20°C; [3]: 1 hour; MS: 791 (M+).

Step 2:

(3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-[4-(3-trifluoromethylbenzyloxy)-benzyl]-3-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carbamate (373)



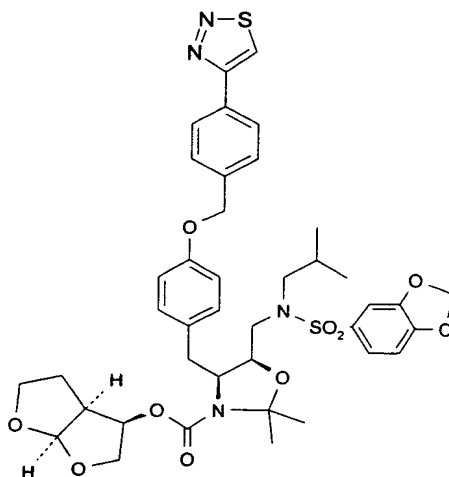
[1]: a, Hexane-EtOAc (1:1); [2]: 47 %. ¹H-NMR (DMSO-d₆): 0.75 (3H,d), 0.81 (3H,d), 1.15-1.22 (1H,m), 1.30 (1H,m), 1.91 (1H,m), 2.34 (1H,t), 2.65-2.75 (3H,m), 2.89-3.01 (2H,m), 3.45 (1H,m), 3.52-3.59 (3H,m), 3.64 (1H,m), 3.80 (1H,dd), 4.80 (1H,dd), 4.99 (1H,d), 5.11 (2H,s(broad)), 5.45 (1H,d), 6.12 (2H,s), 6.84 (2H,d), 7.03 (1H,d), 7.07 (2H,d), 7.21 (2H,m), 7.27 (1H,dd), 7.58 (1H,t), 7.65 (1H,d), 7.70 (1H,d), 7.74 (1H,s); MS: 751(M+).

10

Example 158

Step 1:

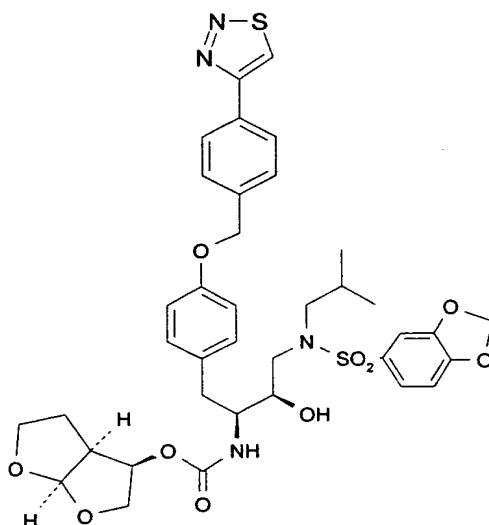
N- (3R,3aS,6aR) -Hexahydrofuro[2,3-b]furan-3-yl-oxycarbonyl-, (4S,5R) -4- [4- (1,2,3-thiadiazole-4-benzyloxy) -benzyl] -5-i-butyl- [(3,4-methylenedioxyphenyl) sulfonyl] -aminomethyl-2,2-dimethyloxazolidine



[1]: 4-[4-(bromomethyl)phenyl]-1,2,3-thiadiazole; [2]:
20°C; [3]: 1 hour; MS: 807 (M+).

Step 2:

- 5 (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-[4-(1,2,3-thiadiazole-4-benzyloxy)-benzyl]-3-i-butyl-[(3,4-methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl)-carbamate (374)

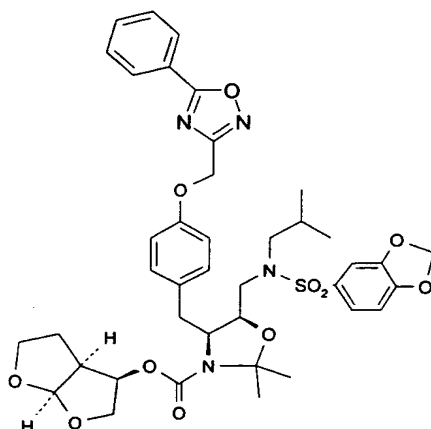


- 10 [1]: a, Hexane-EtOAc (1:3); [2]: 54 %. ¹H NMR (DMSO-d₆):
0.77 (3H,d), 0.83 (3H,d), 1.20 (1H,m), 1.31 (1H,m), 1.93
(1H,m), 2.37 (1H,t), 2.67-2.77 (3H,m), 2.91-3.03 (2H,m),
3.48 (1H,m), 3.53-3.61 (3H,m), 3.67 (1H,m), 3.81 (1H,dd),
4.82 (1H,dd), 5.00 (1H,m), 5.11 (2H,s(broad)), 5.43
15 (1H,d), 6.14 (2H,s), 6.87 (2H,d), 7.04 (1H,d), 7.10
(2H,d), 7.22 (2H,m), 7.29 (1H,d), 7.58 (2H,d), 8.13
(2H,d), 9.59 (1H,s); MS: 767(M+).

Example 159

Step 1:

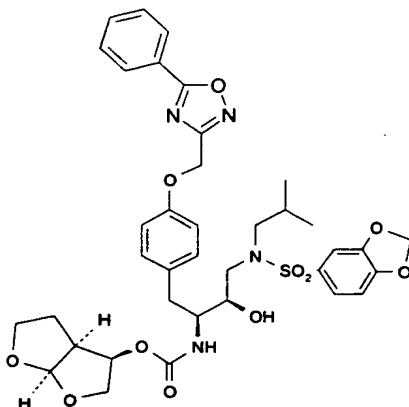
N- (3R,3aS,6aR) -Hexahydrofuro [2,3-b] furan-3-yl-
oxycarbonyl-, (4S,5R) -4- [4- (5-phenyl-1,2,4-oxadiazolo-3-
methyloxy) -benzyl] -5-i-butyl- [(3,4-
5 methylenedioxyphenyl) sulfonyl] -aminomethyl-2,2-dimethyl-
oxazolidine



[1]: 3-chloromethyl-5-phenyl-1,2,4-oxadiazole; [2]: 20°C;
[3]: 24 hour; MS: 791 (M+).

10 Step 2:

(3R,3aS,6aR) -Hexahydrofuro [2,3-b] furan-3-yl-N ((1S,2R) -1-
[4- (5-phenyl-1,2,4-oxadiazolo-3-methyloxy) -benzyl] -3-i-
butyl- [(3,4-methylenedioxyphenyl) sulfonyl] -amino-2-
hydroxypropyl-carbamate (375)



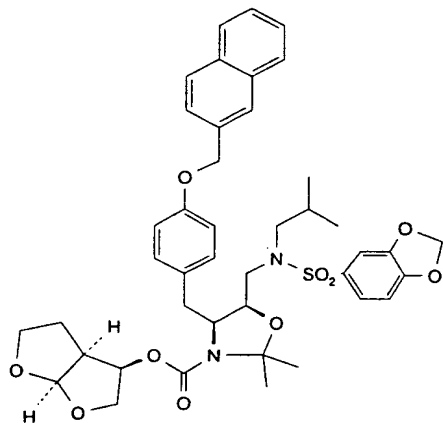
[1]: a, Hexane-EtOAc (1:3); [2]: 53 %. ¹H NMR (DMSO-d₆): 0.77 (3H,d), 0.84 (3H,d), 1.18 (1H,m), 1.32 (1H,m), 1.94 (1H,m), 2.38 (1H,t), 2.68-2.78 (3H,m), 2.93-3.04 (2H,m), 3.46-3.60 (4H,m), 3.69 (1H,m), 3.81 (1H,dd), 4.82 (1H,dd), 5.01 (1H,m), 5.27 (2H,s(broad)), 5.42 (1H,d), 6.14 (2H,s), 6.91 (2H,d), 7.05 (1H,d), 7.12 (2H,d), 7.23 (2H,d), 7.29 (1H,d), 7.62 (2H,m), 7.71 (1H,m), 8.10 (2H,d); MS: 751 (M+).

10

Example 160

Step 1:

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (4S,5R)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 2,2-dimethyl-4-[4-(2-naphthylmethoxy)benzyl]-1,3-
 15 oxazolidine-3-carboxylate



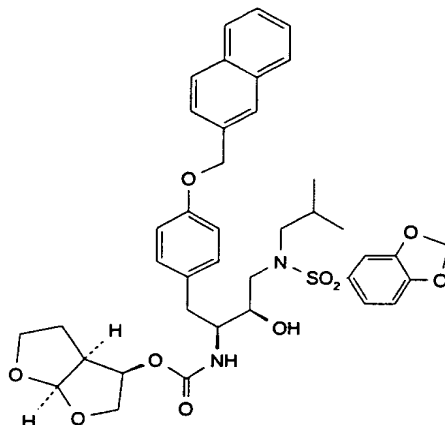
[1]: 2-(bromomethyl)naphthalene; [2]: 20°C; [3]: 12 hour;
 MS: 773 (M+).

20

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-[4-(2-naphthylmethoxy)benzyl]propylcarbamate (376)

5



[1]: c, Et₂O; [2] (Calc. from the product of *Procedure 7*):
 10 92 %.

¹H NMR (DMSO-*d*₆): δ 0.74 (3H,d), 0.80 (3H,d), 1.16 (1H,m),
 1.25 (1H,m), 1.91 (1H,m), 2.33 (1H,t), 2.64-2.73 (3H,m),
 2.88-2.99 (2H,m), 3.44 (1H,m), 3.50-3.56 (3H,m), 3.62
 15 (1H,m), 3.77 (1H,dd), 4.78 (1H,dd), 4.97 (1H, s(broad))
 5.16 (2H,s), 5.40 (1H,d), 6.11 (2H,s), 6.86 (2H,d), 7.01
 (1H,d), 7.06 (2H,d), 7.19 (2H,m), 7.26 (1H,d), 7.46
 (2H,m), 7.50 (1H,d), 7.88 (3H,m); MS: 733 (M+).

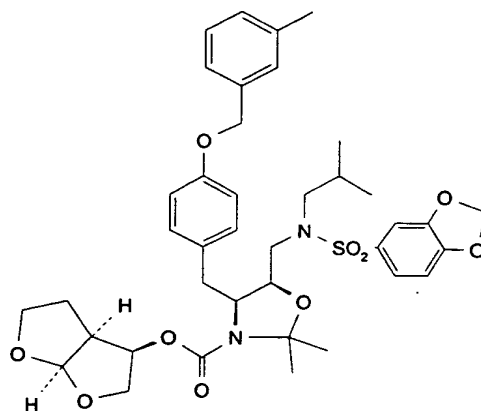
20

Example 161

Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 2,2-dimethyl-4-{4-[(3-methylbenzyl)oxy]benzyl}-1,3-
 oxazolidine-3-carboxylate

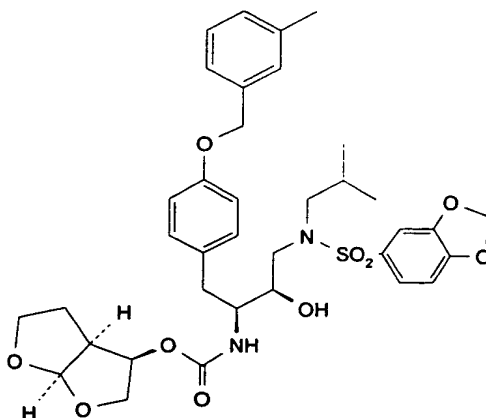
5



[1]: 3-bromomethyltoluene; [2]: 20°C; [3]: 12 hour; MS:
 10 737 (M+).

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy
 15 -1-{4-[(3-methylbenzyl)oxy]benzyl}propylcarbamate (377)



[1]: c, Et₂O; [2]: 72 %.

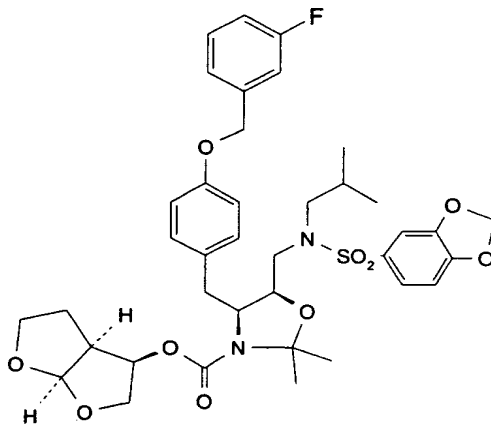
¹H NMR (DMSO-d₆): δ 0.74 (3H,d), 0.80 (3H,d), 1.16 (1H,m),
1.28 (1H,m), 1.90 (1H,m), 2.25 (3H,s), 2.33 (1H,t), 2.64-
5 2.73 (3H,m), 2.88-2.99 (2H,m), 3.44 (1H,m), 3.51-3.57
(3H,m), 3.64 (1H,m), 3.79 (1H,dd), 4.80 (1H,dd), 4.93
(2H,s), 4.97 (1H,s(broad)), 5.45 (1H,d), 6.11 (2H,s),
6.79 (2H,d), 7.00-7.08 (4H,m), 7.14-7.22 (4H,m), 7.25-
7.27 (1H,m); MS: 697 (M+).

10

Example 162

Step 1:

15 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (4S,5R)-5-
{[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
4-{4-[(3-fluorobenzyl)oxy]benzyl}-2,2-dimethyl-1,3-
oxazolidine-3-carboxylate

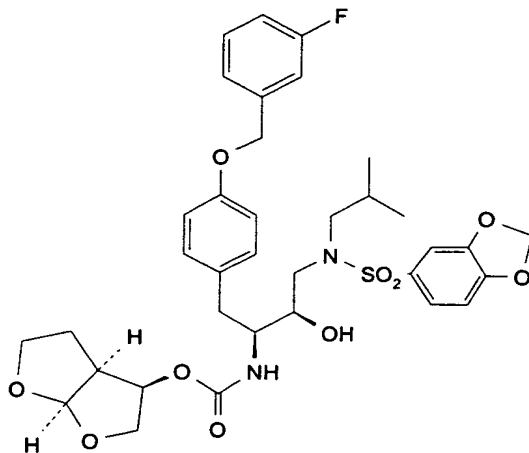


20

[1]: 3-fluorobenzylbromide; [2]: 20°C; [3]: 12 hour; MS:
741 (M+).

Step 2:

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[(3-
 5 fluorobenzyl)oxy]benzyl}-2-hydroxypropylcarbamate (378)



[1]: c, Et₂O; [2]: 67 %.

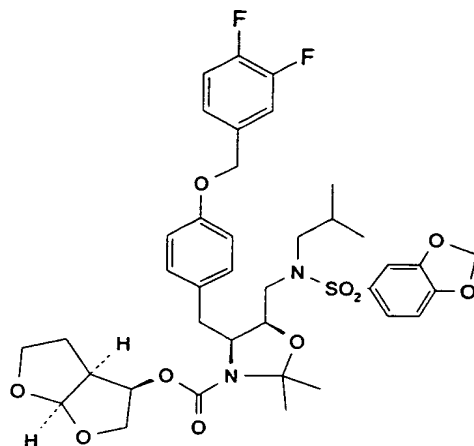
¹H NMR (DMSO-d₆): δ 0.74 (3H,d), 0.80 (3H,d), 1.19 (1H,m),
 10 1.25-1.33 (1H,m), 1.81-1.92 (1H,m), 2.33 (1H,t), 2.64-
 2.73 (3H,m), 2.88-3.00 (2H,m), 3.44 (1H,m), 3.51-3.58
 (3H,m), 3.64 (1H,m), 3.79 (1H,dd), 4.80 (1H,dd), 4.98
 (1H,d) 5.01 (2H,s), 5.45 (1H,d), 6.11 (2H,s), 6.81
 (2H,d), 7.01-7.11 (4H,m), 7.19-7.27 (4H,m), 7.34-7.40
 15 (1H,m); MS: 701 (M⁺).

Example 163

20 Step 1:.

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (4S,5R)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-

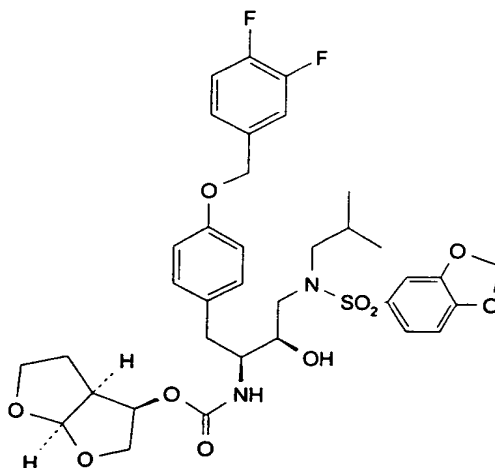
4-{4-[(3,4-difluorobenzyl)oxy]benzyl}-2,2-dimethyl-1,3-oxazolidine-3-carboxylate



5 [1]: α -bromo-3,4-difluorotoluene; [2]: 20°C; [3]: 12 hour; MS: 759 (M+).

Step 2:

10 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[(3,4-difluorobenzyl)oxy]benzyl}-2-hydroxypropylcarbamate (379)



[1]: c, Et₂O; [2]: 65 %.

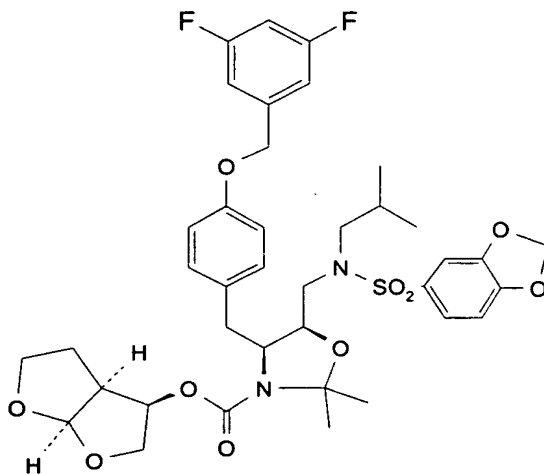
¹H NMR (DMSO-*d*₆): δ 0.74 (3H,d), 0.80 (3H,d), 1.17 (1H,m),
1.27-1.32 (1H,m), 1.88-1.92 (1H,m), 2.33 (1H,t), 2.64-
2.73 (3H,m), 2.88-2.99 (2H,m), 3.42-3.46 (1H,m), 3.53
5 (3H,m), 3.63 (1H,m), 3.78 (1H,dd), 4.80 (1H,dd), 4.97
(3H, s(broad)), 5.45 (1H,d), 6.11 (2H,s), 6.80 (2H,d),
7.00-7.07 (3H,m), 7.19-7.27 (3H,m), 7.35-7.46 (2H,m); MS:
719 (M⁺) .

10

Example 164

Step 1:

15 (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
4-{4-[(3,5-difluorobenzyl)oxy]benzyl}-2,2-dimethyl-1,3-
oxazolidine-3-carboxylate

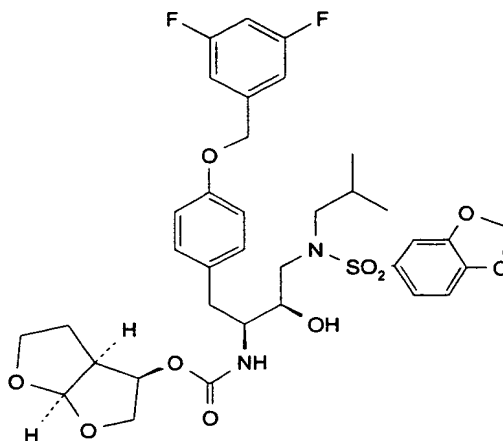


20 [1]: 3,5-difluorobenzylbromide; [2]: 20°C; [3]: 12 hour;
MS: 759 (M⁺) .

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-
 [(3,5-difluorobenzyl)oxy]benzyl}-2-hydroxypropylcarbamate
 (380)

5



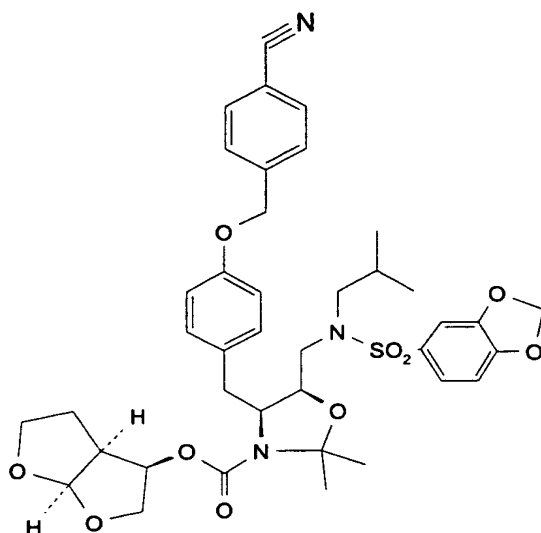
[1]: c, Et₂O; [2]:40 %.

10 ¹H NMR (DMSO-*d*₆): δ 0.74 (3H,d), 0.80 (3H,d), 1.16-1.19
 (1H,m), 1.30 (1H,m), 1.89-1.92 (1H,m), 2.34 (1H,t), 2.64-
 2.73 (3H,m), 2.88-2.99 (2H,m), 3.42-3.46 (1H,m), 3.51-
 3.58 (3H,m), 3.64 (1H,m), 3.79 (1H,dd), 4.80 (1H,dd),
 4.98 (1H,d) 5.03 (2H,s), 5.45 (1H,d), 6.11 (2H,s), 6.81
 15 (2H,d), 7.01-7.15 (6H,m), 7.19-7.27 (2H,m); MS: 719(M⁺).

Example 165

Step 1:

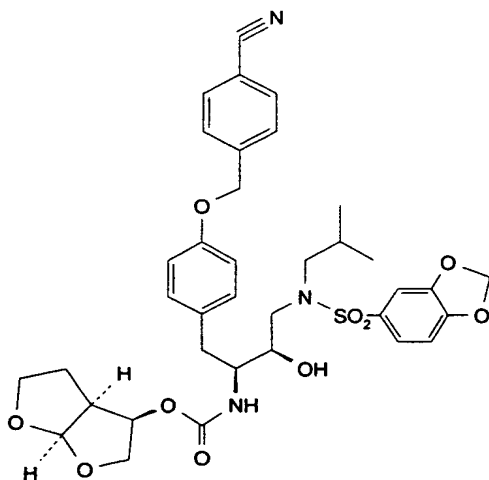
(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
 20 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 4-{4-[(4-cyanobenzyl)oxy]benzyl}-2,2-dimethyl-1,3-
 oxazolidine-3-carboxylate



5 [1]: p-cyanobenzylbromide; [2]: 20°C; [3]: 12 hour; MS:
748 (M+).

Step 2:

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
10 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[(4-
cyanobenzyl)oxy]benzyl}-2-hydroxypropylcarbamate (381)



[1]: b, Hexane-EtOAc (1:3); [2] : 47 %.

^1H NMR ($\text{DMSO}-d_6$): δ 0.75 (3H,d), 0.81 (3H,d), 1.13-1.30 (2H,m), 1.87-1.94 (1H,m), 2.34 (1H,m), 2.62-2.73 (3H,m), 2.89-3.00 (2H,m), 3.43-3.48 (1H,m), 3.51-3.57 (3H,m), 3.61-3.64 (1H,m), 3.79 (1H,dd), 4.80 (1H,dd), 4.99 (1H,d) 5.11 (2H,s), 5.45 (1H,d), 6.12 (2H,s), 6.82 (2H,d), 7.01-7.08 (3H,m), 7.19-7.28 (2H,m), 7.52 (2H,d), 7.81 (2H,d); MS: 708 (M+).

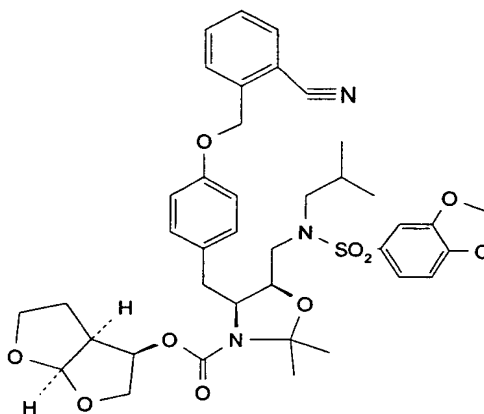
10

Example 166

Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
4-{4-[(2-cyanobenzyl)oxy]benzyl}-2,2-dimethyl-1,3-
oxazolidine-3-carboxylate

15

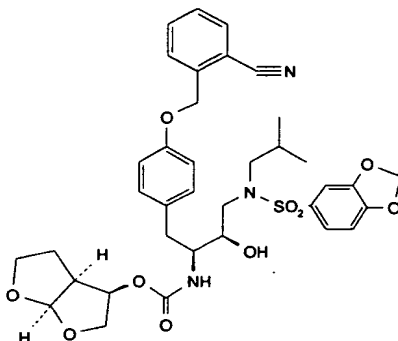


[1]: o-cyanobenzylbromide; [2]: 20°C; [3]: 12 hour; MS:
748 (M+).

20

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[(2-
 cyanobenzyl)oxy]benzyl}-2-hydroxypropylcarbamate (382)



5

[1]: b, Hexane-EtOAc (1:3); [2] :47 %.

1H-NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.19 (1H,m),
 1.31-1.37 (1H,m), 1.91 (1H,m), 2.35 (1H,t), 2.65-2.74
 (3H,m), 2.90-3.00 (2H,m), 3.44-3.49 (1H,m), 3.54 (3H,m),
 10 3.66-3.70 (1H,m), 3.79 (1H,dd), 4.81 (1H,dd), 5.00
 (1H,d), 5.12 (2H,s), 5.45 (1H,d), 6.12 (2H,s), 6.85
 (2H,d), 7.01-7.28 (5H,m), 7.51 (1H,m), 7.62-7.71 (2H,m),
 7.85 (1H,d); MS: 708 (M+).

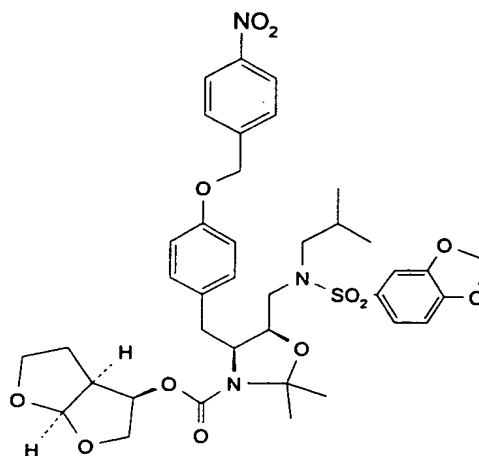
15

Example 167

Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
 20 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 2,2-dimethyl-4-{4-[(4-nitrobenzyl)oxy]benzyl}-1,3-
 oxazolidine-3-carboxylate

25



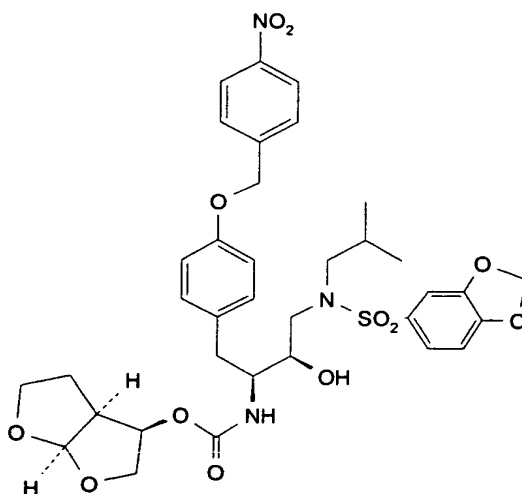
[1]: 4-nitrobenzylbromide; [2]: 20°C; [3]: 3 hour; MS: 768 (M+).

5

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-{4-[(4-nitrobenzyl)oxy]benzyl}propylcarbamate (383)

10



[1]: b, Hexane-EtOAc (1:2); [2]: 50 %.

¹H NMR (DMSO-d₆): δ 0.79 (3H,d), 0.85 (3H,d), 1.24-1.34 (1H,m), 1.95 (1H,m), 2.39 (1H,t), 2.75 (3H,m), 2.93-3.05 (2H,m), 3.34 (1H,m), 3.50-3.68 (5H,m), 3.83 (1H,dd), 4.83 (1H,dd), 5.02 (1H,d), 5.22 (2H,s), 5.48 (1H,d), 6.16 (2H,s), 6.88 (2H,d), 7.05-7.13 (3H,m), 7.24 (2H,m), 7.31 (1H,d), 7.70 (2H,d), 8.24 (2H,d); MS: 728 (M+).

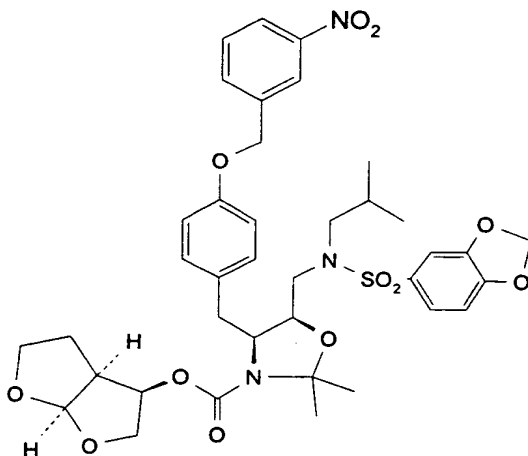
Example 168

Step 1:

10

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (4S,5R)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 2,2-dimethyl-4-{4-[(3-nitrobenzyl)oxy]benzyl}-1,3-
 oxazolidine-3-carboxylate

15

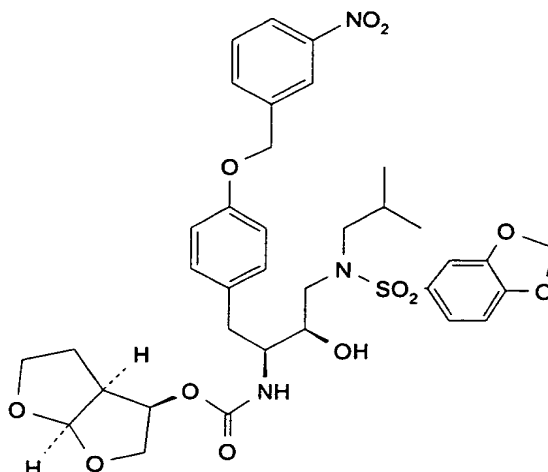


[1]: 3-nitrobenzylbromide; [2]: 20°C; [3]: 3 hour; MS: 768 (M+).

20

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy
 -1-{4-[(3-nitrobenzyl)oxy]benzyl}propylcarbamate (384)



5

[1]: b, Hexane-EtOAc (1:2); [2]: 44 %.

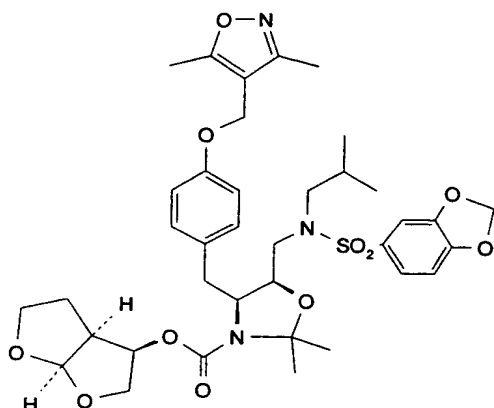
1H-NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.12-1.17
 (1H,m), 1.24-1.30 (1H,m), 1.80-1.94 (1H,m), 2.34 (1H,t),
 2.65-2.74 (3H,m), 2.89-3.00 (2H,m), 3.26 (1H,m), 3.43-3.48
 10 (1H,m), 3.51-3.57 (3H,m), 3.61-3.64 (1H,m), 3.79 (1H,dd),
 4.80 (1H,dd), 4.99 (1H,d), 5.17 (2H,s), 5.44 (1H,d), 6.12
 (2H,s), 6.85 (2H,d), 7.02 (1H,d), 7.08 (2H,d), 7.20-7.22
 (2H,m), 7.26 (1H,dd), 7.64 (1H,t), 7.85 (1H,d), 8.14
 (1H,dd), 8.25 (1H,s); MS: 728 (M+).

15

Example 169

Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
 20 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 4-{4-[(3,5-dimethyl-4-isoxazolyl)methoxy]benzyl}-2,2-
 dimethyl-1,3-oxazolidine-3-carboxylate

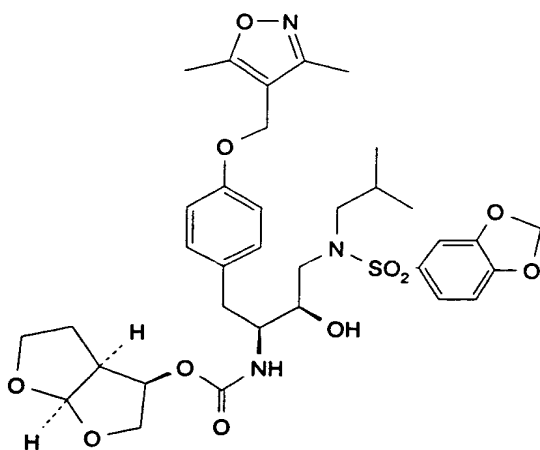


[1]: 4-(chloromethyl)-3,5-dimethylisoxazole; [2]: 20°C;
[3]: 12 hour; MS: 742 (M+).

5

Step 2:

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-
10 [(3,5-dimethyl-4-isoxazolyl)methoxy]benzyl}-2-
hydroxypropylcarbamate (385)



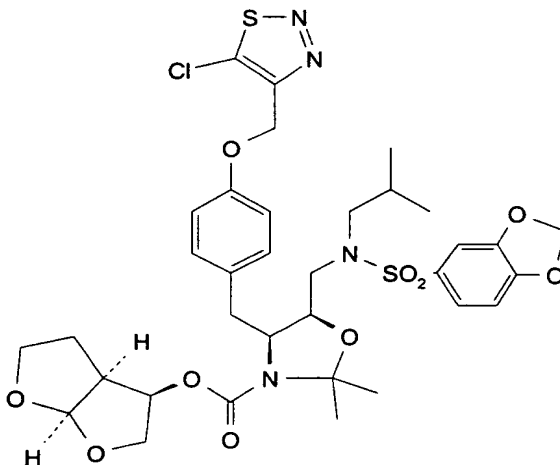
[1]: b, Hexane-EtOAc (1:3); [2] : 49 %.

¹H-NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.19 (1H,m),
1.34 (1H,m), 1.94 (1H,m), 2.14 (3H,s), 2.32 (3H,s), 2.65-
2.74 (3H,m), 2.93 (2H,m), 3.44 (1H,m), 3.52-3.57 (3H,m),
5 3.69 (1H,m), 3.78-3.82 (1H,m), 4.78-4.83 (3H,m), 4.99-
5.00 (1H,m), 5.48 (1H,m), 6.12 (2H,s), 6.81 (2H,d), 7.03
(1H,d), 7.07 (2H,d), 7.20-7.28 (3H,m); MS: 702 (M⁺).

Example 170

10 Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
4-{4-[(5-chloro-1,2,3-thiadiazol-4-yl)methoxy]benzyl}-
15 2,2-dimethyl-1,3-oxazolidine-3-carboxylate

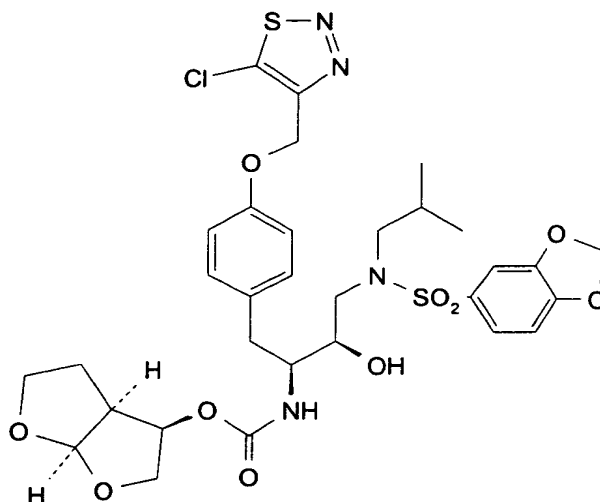


[1]: 5-chloro-4-(chloromethyl)1,2,3-thiadiazole; [2]:
20 20°C; [3]: 12 hour; MS: 765 (M⁺).

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[(5-
chloro-1,2,3-thiadiazol-4-yl)methoxy]benzyl}-2-
hydroxypropylcarbamate (386)

5



[1]: b, Hexane-EtOAc (1:3); [2] : 38 %.

¹H NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.19 (1H,m),
1.32 (1H,m), 1.94 (1H,m), 2.35 (1H,m), 2.67-2.74 (3H,m),
2.91-3.01 (2H,m), 3.46(1H,m), 3.52-3.56 (3H,m), 3.66-3.70
(1H,m), 3.80 (1H,dd), 4.81 (1H,dd), 5.00 (1H,d), 5.35
(2H,s), 5.46 (1H,d), 6.12 (2H,s), 6.90 (2H,d), 7.03
(1H,d), 7.10 (2H,d), 7.20-7.22 (2H,m), 7.26-7.31 (1H,m);
MS: 725 (M+).

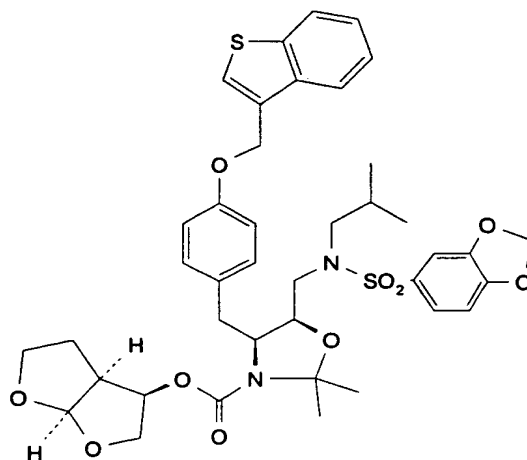
15

Example 171

Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
4-[4-(1-benzothien-3-ylmethoxy)benzyl]-2,2-dimethyl-1,3-
oxazolidine-3-carboxylate

20



[1]: 3-(chloromethyl)-1-benzothiophene; [2]: 20°C; [3]: 12 hour.

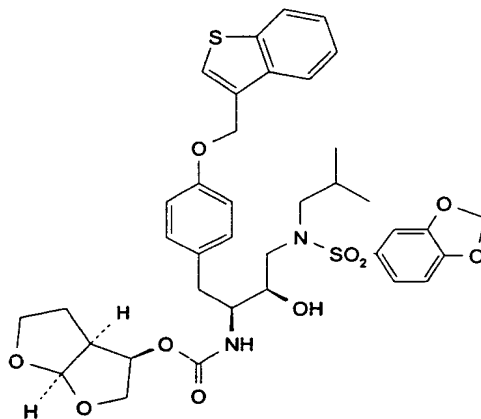
5

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-(1-benzothien-3-ylmethoxy)benzyl]-2-hydroxypropylcarbamate

10

(387)



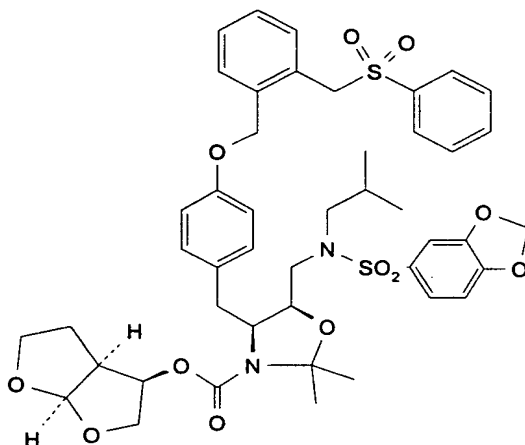
[1]: b, Hexane-EtOAc (1:3); [2]): 8.9 %.

¹H NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.15-1.19 (1H,m), 1.30 (1H,m), 1.90-1.93 (1H,m), 2.35 (1H,t), 2.65-2.74 (3H,m), 2.90-3.00 (2H,m), 3.27 (1H,m), 3.44-3.49 (1H,m), 3.52-3.58 (3H,m), 3.63-3.67 (1H,m), 3.80 (1H,dd), 4.81 (1H,dd), 5.00 (1H,d), 5.24 (2H,s), 5.45 (1H,d), 6.12 (2H,s), 6.89 (2H,d), 7.02 (1H,d), 7.08 (2H,d), 7.20-7.22 (2H,m), 7.26-7.28 (1H,m), 7.33-7.39 (2H,m), 7.80-7.83 (2H,m), 7.95-7.97 (1H,m); MS: 739(M⁺).

10

Example 172

Step 1:



(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 2,2-dimethyl-4-[4-({2-[(phenylsulfonyl)methyl]benzyl}
 oxy)benzyl]-1,3-oxazolidine-3-carboxylate

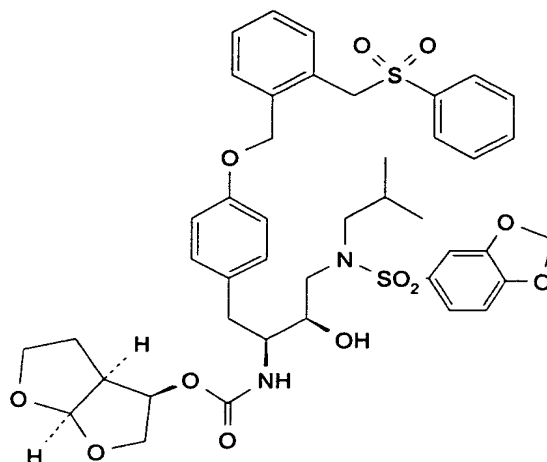
[1]: 1-(bromomethyl)-2-[(phenylsulfonyl)methyl]benzene;
 [2]: 20°C; [3]: 12 hour.

20

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-
 hydroxy-1-[4-({2-[(phenylsulfonyl)methyl]benzyl}oxy)
 benzyl]propylcarbamate (388)

5



[1]: b, Hexane-EtOAc (1:1); [2] :16 %.

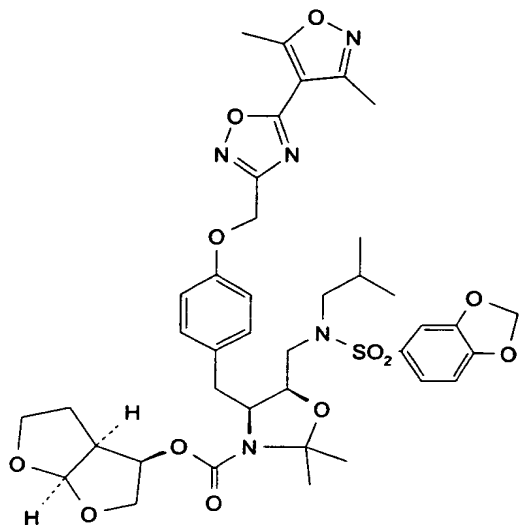
¹H NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.19 (1H,m),
 1.31 (1H,m), 1.90-1.94 (1H,m), 2.35 (1H,t), 2.61-2.74
 10 (3H,m), 2.90-3.01 (2H,m), 3.46 (1H,m), 3.52-3.56 (3H,m),
 3.66 (1H,m), 3.79 (1H,dd), 4.75 (2H,s), 4.79-4.84 (1H,m),
 4.99-5.02 (3H,m), 5.45 (1H,d), 6.12 (2H,s), 6.79 (2H,d),
 7.01-7.08 (4H,m), 7.02 (2H,m), 7.23-7.32 (3H,m), 7.40
 (1H,d), 7.54-7.61 (2H,m), 7.68-7.76 (3H,m); MS: 837(M⁺).

15

Example 173

Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 20 4-(4-{[5-(3,5-dimethyl-4-isoxazolyl)-1,2,4-oxadiazol-3-
 yl]methoxy}benzyl)-2,2-dimethyl-1,3-oxazolidine-3-
 carboxylate

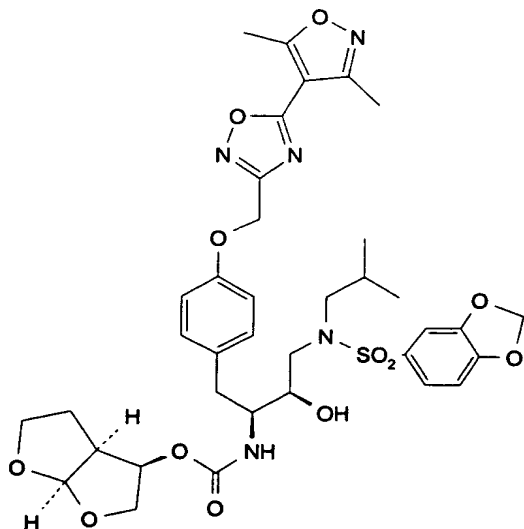


[1]: 3-(chloromethyl)-5-(3,5-dimethyl-4-isoxazolyl)-
 5 1,2,4-oxadiazole; [2]: 20°C; [3]: 1 hour.

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-(4-{[5-
 10 (3,5-dimethyl-4-isoxazolyl)-1,2,4-oxadiazol-3-
 yl]methoxy}benzyl)-2-hydroxypropylcarbamate (389)

[1]: b, Hexane-EtOAc (1:3); [2]: 4 %.



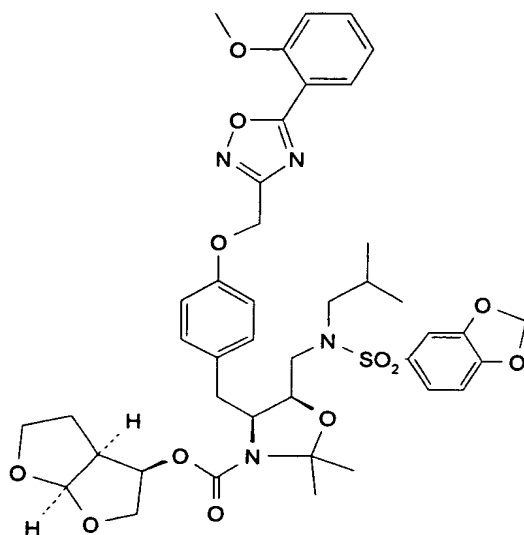
¹H NMR (DMSO-*d*₆): δ 0.74 (3H,d), 0.81 (3H,d), 1.13-1.28 (2H,m), 1.90 (1H,m), 2.31-2.37 (2H,m), 2.71 (6H,m), 2.90-3.00 (2H,m), 3.46-3.53 (4H,m), 3.64 (1H,m), 3.78 (1H,m), 4.80 (1H,m), 5.01 (1H,m), 5.23 (2H,m), 6.11 (2H,s), 6.88 (2H,d), 7.02 (1H,dd), 7.09 (1H,d), 7.19-7.22 (2H,m), 7.26 (1H,d); MS: 770 (M+).

Example 174

Step 1:

10

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
4-(4-{[5-(2-methoxyphenyl)-1,2,4-oxadiazol-3-
yl]methoxy}benzyl)-2,2-dimethyl-1,3-oxazolidine-3-
15 carboxylate



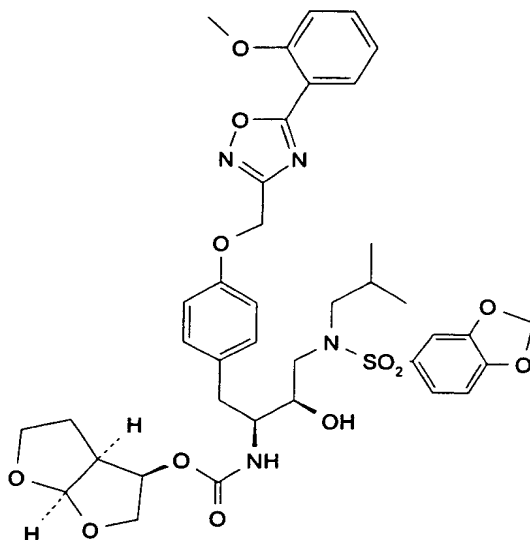
[1]: 3-(chloromethyl)-5-(2-methoxyphenyl)-1,2,4-oxadiazole; [2]: 20°C; [3]: 1 hour.

20

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-{[5-(2-methoxyphenyl)-1,2,4-oxadiazol-3-yl]methoxy}benzyl)propylcarbamate (390)

5



[1]: b, Hexane-EtOAc (1:2); [2]:24 %.

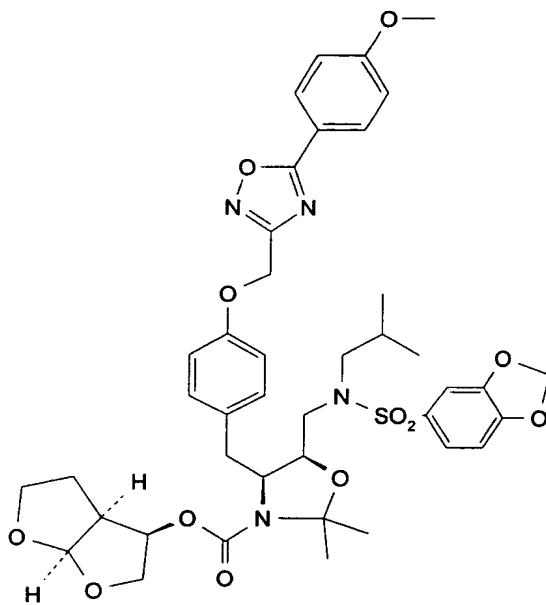
¹H NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.12-1.19 (1H,m), 1.26-1.37 (1H,m), 1.90-1.93 (1H,m), 2.28-2.38 (1H,m), 2.62-2.74 (3H,m), 2.90-3.00 (2H,m), 3.45 (1H,m), 3.51-3.55 (3H,m), 3.64-3.69 (1H,m), 3.78 (1H,dd), 3.88 (3H,s), 4.79 (1H,dd), 5.00 (1H,d), 5.22 (2H,s), 5.40 (1H,d), 6.12 (2H,s), 6.88 (2H,d), 7.02 (1H,d), 7.08-7.12 (3H,m), 7.20-7.23 (2H,m), 7.26 (2H,d), 7.63 (1H,t), 7.94 (1H,d); MS: 781 (M⁺).

Example 175

20 Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-[[[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl]-

4-(4-{[5-(4-methoxyphenyl)-1,2,4-oxadiazol-3-yl]methoxy}benzyl)-2,2-dimethyl-1,3-oxazolidine-3-carboxylate



5

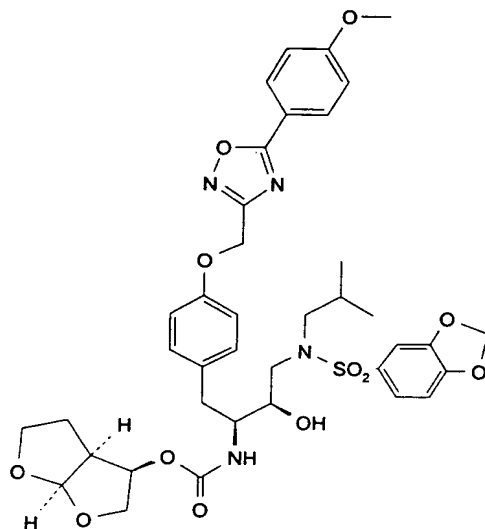
[1]: 3-(chloromethyl)-5-(4-methoxyphenyl)-1,2,4-oxadiazole; [2]: 20°C; [3]: 1 hour.

10 Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-2-hydroxy-1-(4-{[5-(4-methoxyphenyl)-1,2,4-oxadiazol-3-yl]methoxy}benzyl)propylcarbamate (391)

15

20



5

[1]: a, Hexane-EtOAc (1:5); [2] : 44 %.

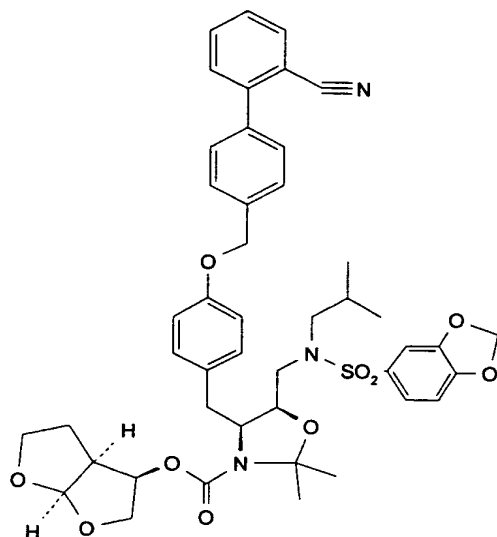
¹H NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.12-1.19 (1H,m), 1.24-1.36 (1H,m), 1.91 (1H,m), 2.32-2.38 (1H,m), 2.62-2.74 (3H,m), 2.90-3.00 (2H,m), 3.46 (1H,m), 3.51-3.55 (3H,m), 3.66 (1H,m), 3.76-3.79 (1H,m), 3.82 (3H,m), 4.79 (1H,dd), 5.00 (1H,d), 5.20 (2H,s), 5.40 (1H,d), 6.12 (2H,s), 6.88 (2H,d), 7.02 (1H,d), 7.08-7.13 (3H,m), 7.20-7.23 (2H,m), 7.26 (1H,d), 8.02 (2H,d); MS: 781 (M⁺).

15

Example 176

Step 1:

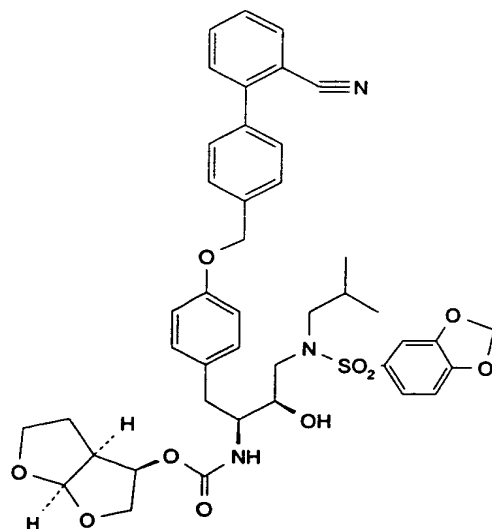
(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 4-{4-[(2'-cyano[1,1'-biphenyl]-4-yl)methoxy]benzyl}-2,2-
 dimethyl-1,3-oxazolidine-3-carboxylate



[1]: 4'-(bromomethyl)[1,1'-biphenyl]-2-carbonitrile; [2]:
 5 20°C; [3]: 3 hour; MS: 824 (M+).

Step 2:

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-
 10 [(2'-cyano[1,1'-biphenyl]-4-yl)methoxy]benzyl}-2-
 hydroxypropylcarbamate (392)



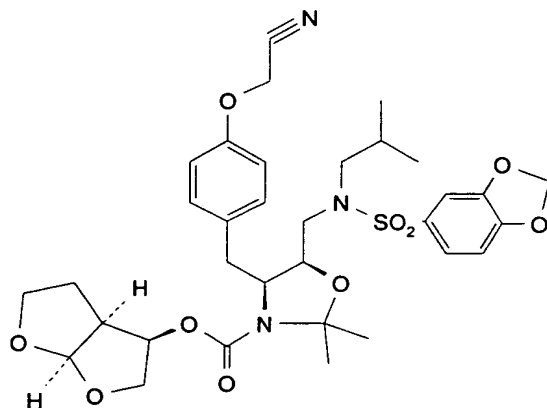
[1]: b, Hexane-EtOAc (1:2); [2] :54 %.

¹H NMR (DMSO-d₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.19 (1H,m),
1.26-1.38 (1H,m), 1.92 (1H,m), 2.32-2.38 (1H,m), 2.65-
5 2.75 (3H,m), 2.90-3.01 (2H,m), 3.30 (1H,m), 3.46 (1H,m),
3.52-3.56 (3H,m), 3.66 (1H,m), 3.79 (1H,dd), 4.81
(1H,dd), 4.99 (1H,d), 5.08 (2H,s), 5.44 (1H,d), 6.12
(2H,s), 6.86 (2H,d), 7.02 (1H,d), 7.08 (2H,d), 7.20-7.22
(2H,m), 7.27 (1H,dd), 7.52-7.60 (6H,m), 7.75 (1H,t), 7.91
10 (1H,d); MS: 784 (M+).

Example 177

Step 1:

15 (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
4-[4-(cyanomethoxy)benzyl]-2,2-dimethyl-1,3-oxazolidine-
3-carboxylate

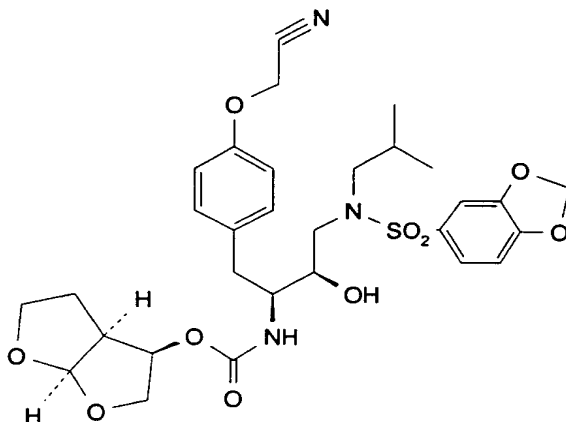


20

[1]: chloroacetonitrile; [2]: 20°C; [3]: 3 hour; MS: 672
(M+).

Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-[4-
 (cyanomethoxy)benzyl]-2-hydroxypropylcarbamate (393)



5

[1]: c, Hexane-EtOAc (50:1); [2] : 48 %.

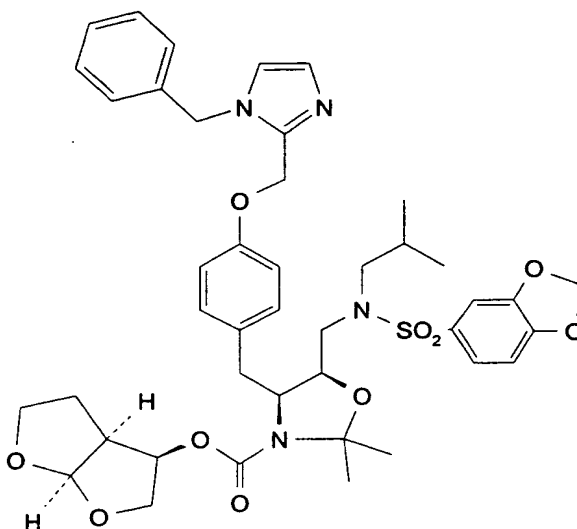
¹H NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.20 (1H,m),
 1.36-1.41 (1H,m), 1.91 (1H,m), 2.31-2.37 (1H,m), 2.65-
 10 2.74 (3H,m), 2.89-3.00 (2H,m), 3.43(1H,m), 3.52-3.60
 (2H,m), 3.71 (1H,m), 3.78-3.82 (1H,m), 4.29-4.34 (2H,m),
 4.79-4.84 (1H,m), 5.00 (1H,d), 5.46 (1H,d), 6.12 (2H,s),
 6.76 (2H,d), 7.01-7.07 (3H,m), 7.20-7.28 (1H,m), 7.26
 (1H,m), 7.33 (1H,m), 7.45 (1H,m); MS: 632 (M+).

15

Example 178

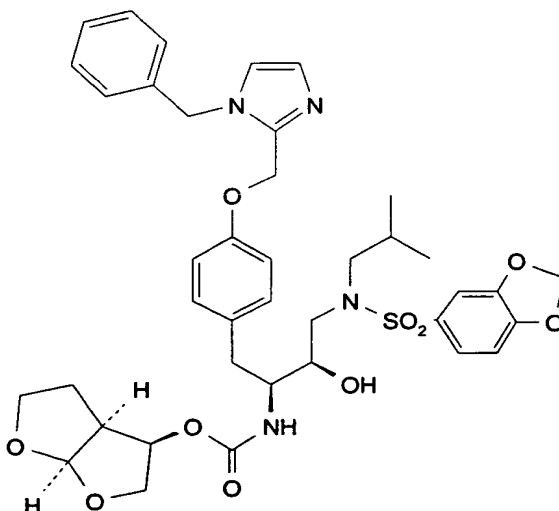
Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
 20 {[(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]methyl}-
 4-{4-[(1-benzyl-1*H*-imidazol-2-yl)methoxy]benzyl}-2,2-
 dimethyl-1,3-oxazolidine-3-carboxylate



[1]: 1-benzyl-2-(chloromethyl)-1H-imidazole
 5 hydrochloride; [2]: 20°C; [3]: 3 hour; MS: 803 (M+).

Step 2:
 (3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(isobutyl)amino]-1-{4-[(1-
 10 benzyl-1H-imidazol-2-yl)methoxy]benzyl}-2-
 hydroxypropylcarbamate (394)



[1]: b, EtOAc-EtOH(1:1); [2] : 29 %.

¹H-NMR (DMSO-*d*₆): δ 0.75 (3H,d), 0.81 (3H,d), 1.11-1.32 (2H,m), 1.88-1.94 (1H,m), 2.28-2.40 (1H,m), 2.65-2.74 (3H,m), 2.86-3.01 (2H,m), 3.26(1H,m), 3.43-3.48 (1H,m), 3.52-3.57 (2H,m), 3.62-3.66 (1H,m), 3.79 (1H,dd), 4.81 (1H,dd), 4.95-5.05 (3H,m), 5.19 (2H,s), 5.45 (1H,d), 6.12 (2H,s), 6.80 (2H,d), 6.88 (1H,m), 7.02-7.06 (3H,m), 7.11-7.15 (2H,m), 7.20-7.30 (6H,m); MS: 763 (M+).

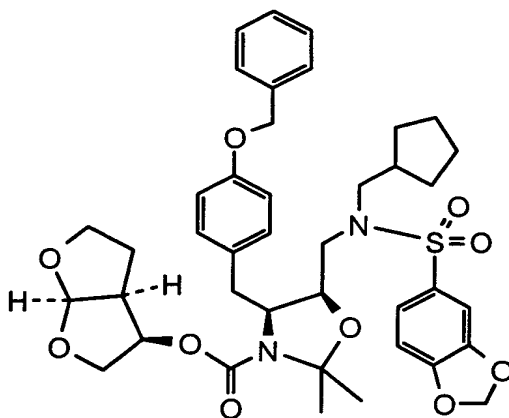
10

Example 179

Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[(1,3-benzodioxol-5-ylsulfonyl)(cyclopentylmethyl)
amino]methyl}-4-[4-(benzyloxy)benzyl]-2,2-dimethyl-1,3-
oxazolidine-3-carboxylate

15



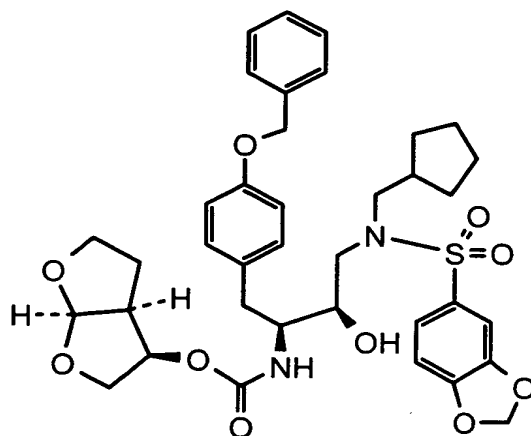
The reaction was carried out as described for N-
(3*R*,3*aS*,6*aR*)-Hexahydrofuro[2,3-*b*]furan-3-yl-oxycarbonyl-,
(4*S*,5*R*)-4-(4-benzyloxy-benzyl)-5-*i*-butyl-[(3,4-
methylenedioxyphenyl)sulfonyl]-aminomethyl-2,2-dimethyl-
oxazolidine. (Y=67%)

20

¹H NMR (CDCl₃): δ 1.08 (1H,m), 1.25 (2H,m), 1.45-1.80 (5H,m), 1.48 (3H,s), 1.64 (3H,s), 1.85 (2H,m), 2.25 (1H,m), 2.65-3.45 (7H,m), 3.80 (3H,m), 3.95 (1H,m), 4.21 (1H,m), 4.28 (1H,m), 4.88 (1H,dd), 5.03 (2H, s), 5.65 (1H,d), 6.01 (2H,s), 6.80-7.50 (12H,m).

Step 2:

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 10 [(1,3-benzodioxol-5-ylsulfonyl)(cyclopentylmethyl)amino]-
 1-[4-(benzyloxy)benzyl]-2-hydroxypropylcarbamate (395)



15 The carbamate formation was carried out as described for
 (3R,3aS,6aR)-Hexahydrofuro[2,3-b]furan-3-yl-N((1S,2R)-1-
 (4-benzyloxy-benzyl)-3-i-butyl-[(3,4-
 methylenedioxyphenyl)sulfonyl]-amino-2-hydroxypropyl-
 caramate. (Y=79%)

20

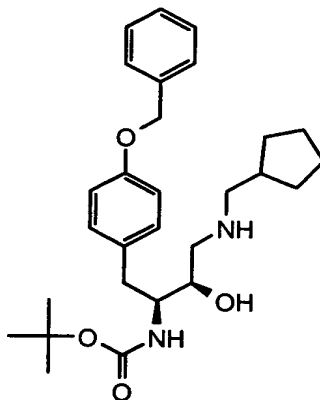
¹H NMR (DMSO-d₆): δ 1.05 (1H,m), 1.20 (1H,m), 1.30-1.70 (8H,m), 2.19 (1H,m), 2.36 (1H,t), 2.70-2.95 (4H,m), 3.10 (1H,dd), 3.40-3.60 (4H,m), 3.65 (1H,t), 3.80 (1H,dd),

4.81 (1H,dd), 5.00 (3H, s+d), 5.46 (1H,d), 6.13 (2H,s),
6.83 (2H,d), 7.00-7.41 (11H,m). MS: 708 (M)+.

Example 180

5 Step 1:

**tert-butyl (1*S*,2*R*)-1-[4-(benzyloxy)benzyl]-3-
[(cyclopentylmethyl)amino]-2-hydroxypropylcarbamate**



10

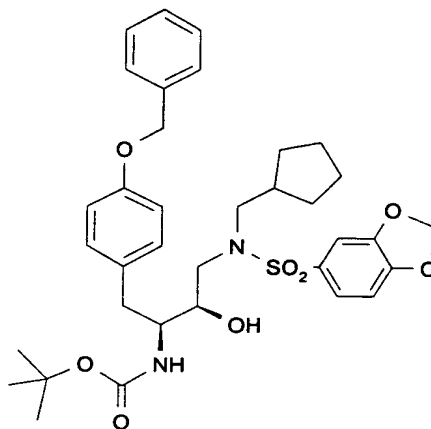
The reaction was carried out as described above except
cyclopentylmethylamine was used in the reaction. (Y:
91%)

15 ¹H NMR: (CDCl₃): δ 1.08-1.23 (2H,m), 1.34 (9H,s), 1.48-
1.59 (4H,m), 1.72-1.77 (2H,m), 1.91-2.02 (1H,m), 2.45-
2.54 (2H,m), 2.67 (2H,m), 2.77-2.82 (1H,m), 2.88 (1H,dd),
3.41 (1H,m), 3.73 (1H,m), 4.64 (1H,d), 5.01 (2H,s), 6.88
20 (2H,d), 7.12 (2H,d), 7.28-7.41 (4H,m).

20

Step 2:

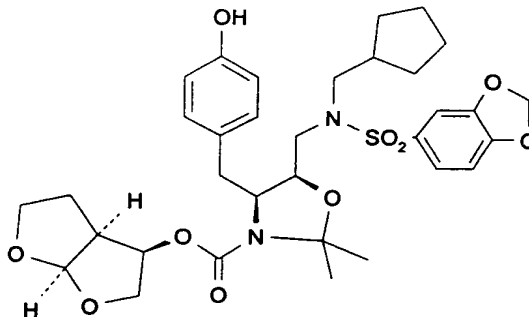
tert-butyl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)
(cyclopentylmethyl) amino]-1-[4-(benzyloxy)benzyl]-2-
hydroxypropylcarbamate



- 5 The reaction was carried out as described above except
tert-butyl (1*S*,2*R*)-1-[4-(benzyloxy)benzyl]-3-
[(cyclopentylmethyl) amino]-2-hydroxypropylcarbamate was
used in the reaction. (Y: 88%)
This compound was used as is in the next step without
10 additional purification.

Step 3:

- (3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[(1,3-benzodioxol-5-ylsulfonyl)(cyclopentylmethyl)
15 amino]methyl}-4-(4-hydroxybenzyl)-2,2-dimethyl-1,3-
oxazolidine-3-carboxylate



This reaction was carried out as described above except
tert-butyl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-
ylsulfonyl)(cyclopentylmethyl)amino]-1-[4-
(benzyloxy)benzyl]-2-hydroxypropylcarbamate was used in
5 the reaction. (Y: 59%)

¹H NMR: (CDCl₃): δ 1.02-1.12 (1H,m), 1.20-1.33 (1H,m),
1.45-1.76 (5H,m), 1.47 (3H,s), 1.65 (3H,s), 1.87 (2H,m),
2.16-2.30 (1H,m), 2.66-2.70 (2H,m), 2.78-2.88 (2H,m),
10 2.98-3.04 (1H,m), 3.11-3.20 (1H,m), 3.37-3.41 (2H,m),
3.75-3.85 (2H,m), 3.91-3.96 (1H,m), 4.17-4.21 (1H,m),
4.26-4.28 (1H,m), 4.87 (1H,dd), 5.66 (1H,dd), 6.04
(2H,s), 6.74 (2H,d), 6.82 (1H,d), 6.96 (2H,d), 7.02
(1H,d), 7.04-7.07 (1H,m), 7.11 (1H,dd)

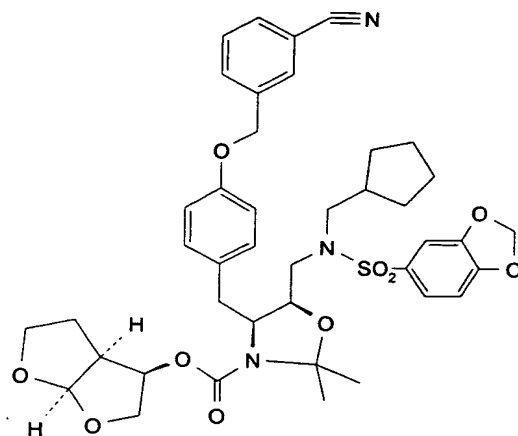
15

Step 4:

General procedure for the aralkylation of (3*R*,3*aS*,6*aR*)-
hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-{[(1,3-
benzodioxol-5-ylsulfonyl)(cyclopentylmethyl)
20 amino]methyl}-4-(4-hydroxybenzyl)-2,2-dimethyl-1,3-
oxazolidine-3-carboxylate

This procedure was carried out as described before using
aralkyl halide [1], stirring at the indicated temperature
25 [2] for the indicated number of hours [3].

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
{[(1,3-benzodioxol-5-ylsulfonyl)(cyclopentylmethyl)
30 amino]methyl}-4-{4-[(3-cyanobenzyl)oxy]benzyl}-2,2-
dimethyl-1,3-oxazolidine-3-carboxylate



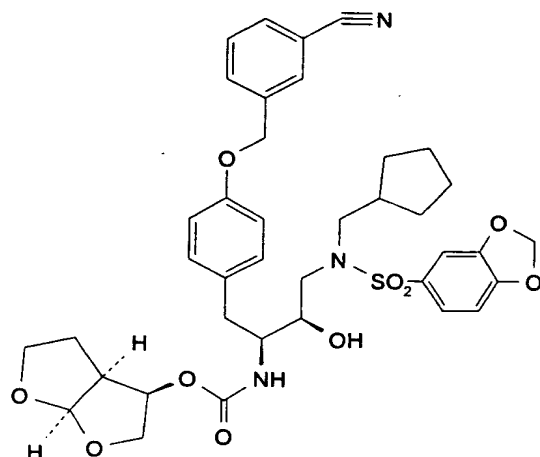
[1]: 3-cyanobenzyl bromide; [2]: 20°C; [3]: 12 hour; MS:
5 774 (M+).

Step 5:

General Procedure for the deprotection

10 This reaction was carried out as before using HCl/dioxane

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-
[(1,3-benzodioxol-5-ylsulfonyl)(cyclopentylmethyl)amino]-
1-{4-[(3-cyanobenzyl)oxy]benzyl}-2-hydroxypropylcarbamate
15 (396)



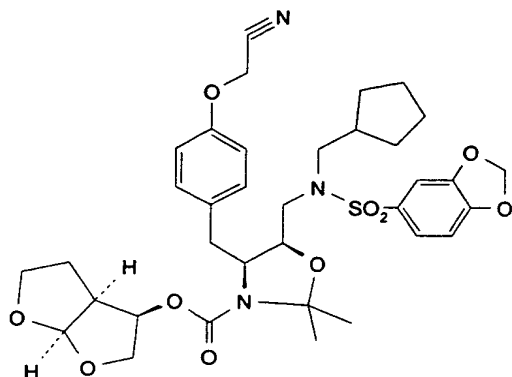
[1]: c, EtOAc-Hexane; [2] :60%.

5 1H-NMR: (DMSO): δ 1.10 (1H,m), 1.17-1.32 (2H,m), 1.35-1.72 (5H,m), 2.21 (1H,m), 2.39 (1H,m), 2.74-3.00 (4H,m), 3.11-3.18 (1H,m), 3.46-3.62 (4H,m), 3.64-3.70 (1H,m), 3.84 (1H,dd), 4.85 (1H,dd), 5.03 (1H,m), 5.11 (2H,s), 5.50 (1H,d) 6.16 (2H,s), 6.87 (2H,d), 7.05-7.13 (3H,m),
 10 7.25 (2H,d), 7.31 (1H,d), 7.60 (1H,t), 7.76-7.80 (2H,m), 7.88 (1H,s); MS: 734 (M+).

Example 181

15 Step 1

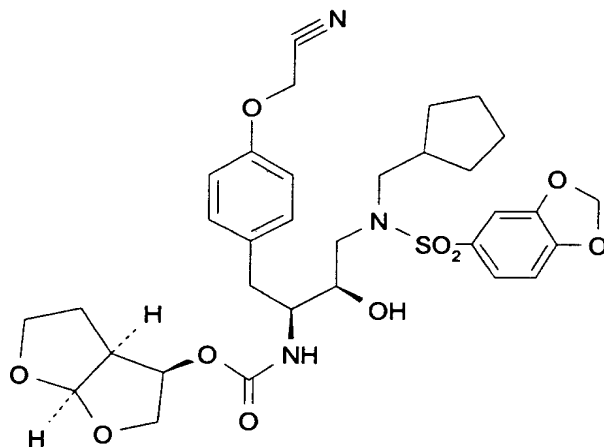
(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (4S,5R)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(cyclopentylmethyl)
 amino]methyl}-4-[4-(cyanomethoxy)benzyl]-2,2-dimethyl-
 1,3-oxazolidine-3-carboxylate



[1]: chloroacetonitrile; [2]: 20°C; [3]: 12 hour; MS: 698 (M+).

5 Step 2:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (1*S*,2*R*)-3-[(1,3-benzodioxol-5-ylsulfonyl)(cyclopentylmethyl)amino]-1-[4-(cyanomethoxy)benzyl]-2-hydroxypropylcarbamate (397)



10

[1]: b, EtOAc-Hexane (2:1); [2] : 77%.

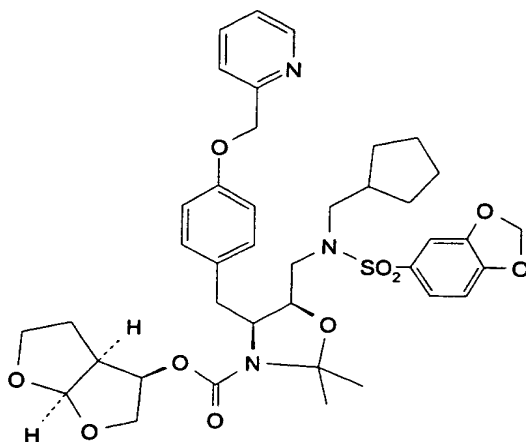
¹H NMR: (DMSO): δ 1.10 (1H,m), 1.24 (2H,m), 1.40-1.70 (7H,m), 2.20-2.26 (1H,m), 2.35-2.43 (1H,m), 2.78-2.98 (4H,m), 3.11-3.19 (1H,m), 3.48-3.51 (1H,m), 3.55-3.65 (3H,m), 3.73-3.78 (1H,m), 3.84 (1H,dd), 4.33 (2H,s), 4.85 (1H,dd), 5.02 (1H,d), 5.51 (1H,d), 6.16 (2H,s), 6.80 (2H,d), 7.05-7.12 (3H,m), 7.22-7.25 (1H,m), 7.31 (1H,dd), 7.36 (1H,m), 7.46 (1H,m); MS: 658 (M+).

10

Example 182

Step 1:

(3*R*,3*aS*,6*aR*)-hexahydrofuro[2,3-*b*]furan-3-yl (4*S*,5*R*)-5-
 {[(1,3-benzodioxol-5-ylsulfonyl)(cyclopentylmethyl)
 15 amino]methyl}-2,2-dimethyl-4-[4-(2-
 pyridinylmethoxy)benzyl]-1,3-oxazolidine-3-carboxylate

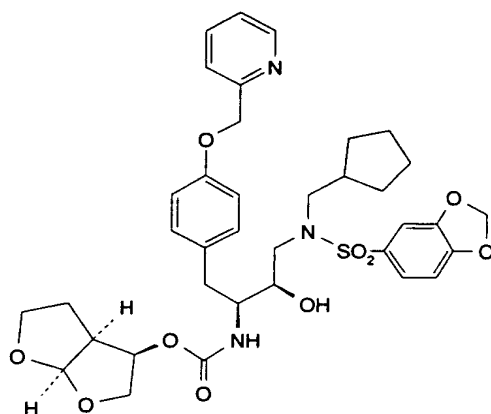


20 [1]: 2-picolychloride HCl; [2]: 20°C; [3]: 12 hour; MS:
 750 (M+).

Step 2:

(3R,3aS,6aR)-hexahydrofuro[2,3-b]furan-3-yl (1S,2R)-3-
 [(1,3-benzodioxol-5-ylsulfonyl)(cyclopentylmethyl)amino]-
 2-hydroxy-1-[4-(2-pyridinylmethoxy)benzyl]propylcarbamate
 (398)

5



[1]: b, EtOAc; [2] :47%.

10 ¹H NMR: (DMSO): δ 1.14 (1H,m), 1.24 (2H,m), 1.40-1.70
 (7H,m), 2.22 (1H,m), 2.39 (1H,m), 2.74-2.96 (4H,m), 3.10-
 3.15 (1H,m), 3.51-3.59 (4H,m), 3.63-3.70 (1H,m), 3.83
 (1H,dd), 4.84 (1H,dd), 5.01 (1H,m), 5.10 (2H,s), 5.27
 (1H,d) 6.15 (2H,s), 6.87 (2H,d), 7.04-7.12 (3H,m), 7.21-
 15 7.25 (2H,m), 7.30-7.34 (2H,m), 7.48 (1H,d), 7.81 (1H,t),
 8.55 (1H,d); MS: 710 (M+).

Example 183

Anti-Viral Activity

20 We measured the enzyme inhibition constants of
 the compounds listed in Table I against HIV-1 protease
 using the methods of: B. Maschera et al., "Human
 Immunodeficiency Virus: Mutations in the Viral Protease
 that Confer Resistance to Saquinavir Increase the

Dissociation Rate Constant for the Protease-Saquinavir Complex", J. Biol. Chem., 271, pp. 33231-35 (1996); and M. V. Toth et al., Int. J. Peptide Protein Res. 36, pp. 544-50 (1990)

5

Antiviral activity assay in MT4 cells

Antiviral HIV activity and compound-induced cytotoxicity were measured in parallel by means of a propidium iodide based procedure in the human T-cell lymphotropic virus transformed cell line MT4. Aliquots of the test compounds were serially diluted in medium (RPMI 1640, 10% fetal calf serum (FCS), and gentamycin) in 96-well plates (Costar 3598) using a Cetus Pro/Pette. Exponentially growing MT4 cells were harvested and centrifuged at 1000 rpm for 10 min in a Jouan centrifuge (model CR 4 12). Cell pellets were resuspended in fresh medium (RPMI 1640, 20% FCS, 20% IL-2, and gentamycin) to a density of 5×10^5 cells/ml. Cell aliquots were infected by the addition of HIV-1 (strain IIIB) diluted to give a viral multiplicity of infection of 100 x TCID₅₀. A similar cell aliquot was diluted with medium to provide a mock-infected control. Cell infection was allowed to proceed for 1 hr at 37° in a tissue culture incubator with humidified 5% CO₂ atmosphere. After the 1 hr incubation the virus/cell suspensions were diluted 6-fold with fresh medium, and 125 µl of the cell suspension was added to each well of the plate containing pre-diluted compound. Plates were then placed in a tissue culture incubator with humidified 5% CO₂ for 5 days. At the end of the incubation period, 27 µl of 5% Nonidet-40 was added to each well of the incubation plate. After thorough mixing with a Costar multitip pipetter, 60 µl of

the mixture was transferred to filter-bottomed 96-well plates. The plates were analyzed in an automated assay instrument (Screen Machine, Idexx Laboratories). The assay makes use of a propidium iodide dye to estimate the DNA content of each well.

REFERENCES

1. Averett, D.R. 1989. Anti-HIV compound assessment by two novel high capacity assays. *J. Virol. Methods* 23: 263-276.
2. Schwartz, O., et al. 1988. A rapid and simple colorimetric test for the study of anti-HIV agents. *AIDS Res. and Human Retroviruses*, 4(6):441-447.
3. Daluge, S.M., et al. 1994. 5-chloro-2'3'-deoxy-3'fluorouridine (935U83), a selective anti-human immuno-deficiency virus agent with an improved metabolic and toxicological profile. *Antimicro. Agents and Chemother.*, 38 (7):1590-1603.

The anti-viral potency in MT-4 cells of the compounds set forth in Tables 1 and 2 was determined using the above technique. The results are shown in Table A.

Table A. Antiviral Activity of Compounds of the Invention.

Cmpd #	IC ₅₀	Cmpd #	IC ₅₀	Cmpd #	IC ₅₀
5a	NA	26	A	48	B
5b	NA	27	D	49	B
5c	NA	28	E	50	C

5d	NA	29	C	51	B
5e	NA	30	C	52	B
5f	NA	31	B	53	B
5g	NA	32	C	54	B
6e	NA	33	C	55	A
10	D	34	D	56	C
11	D	35	C	57	B
12	E	36	C	58	E
13	D	37	C	59	NA
14	NA	38	A	60	NA
15	D	39	NA	61	NA
16	D	40	NA	67	D
17	D	41	NA	68	D
18	C	42	D	69	D
19	E	43	C	70	D
20	C	44	A	71	B
21	C	46	A	72	B
22	C	47	NA	73	C
23	C			74	D
24	C				
25	C				

In Table A, the following classifications have been employed:

A < 0.001 μ M

0.010 > B > 0.001 μ M

0.100 > C > 0.010 μ M

D > 0.1 μ M

"NA" = compound was not tested.

5

Anti-viral activity against Resistant Viruses

EP13 and D545701, two multi protease inhibitor resistant viruses were used to assess potency against mutant viruses. These viruses contain the following
10 mutations relative to the consensus sequence of wild type virus:

D545701-14: Protease amino acid sequence: L10I, L19Q, K20R, E35D, M36I, S37N, M46I, I50V, I54V, I62V, L63P,
15 A71V, V82A, L90M; reverse transcriptase amino acid sequence: E28K, K32E, V35I, T39S/T, E40D/V/Y/F, M41M/L, K43E, Y181Y/C

EP13: Protease amino acid sequence: M46I, L63P, A71V,
20 V82F/L, I84V; No reverse transcriptase mutations.

Reference data for the following protease inhibitors are (D545701-14; EP13):

Amprenavir™: >1000nM; 600nM
25 Indinavir™: 700nM; 560nM
Nelfinavir™: 690nM; N/A
Ritonavir™: >1000nM; >600nM
Saquinavir™: 900nM; N/A

30 Assays of the above mutant viruses were carried out as described above for the wild type virus and the results are shown below in Table B.

Table B

Compound No.	Wildtype virus - IC_{50}	EP13 mutant IC_{50}	D545701-14 mutant - IC_{50}
52	C	B	B
53	B	B	B
54	C	B	B
55	NA	B	B
201	NA	NA	NA
202	A	A	B
203	A	B	C
204	B	B	B
205	B	C	C
206	B	C	C
207	B	B	B
208	C	NA	NA
209	B	B	B
210	B	B	C
211	B	B	B
212	B	B	C
213	B	B	B
214	B	B	B
215	B	B	B
216	B	B	B
217	B	B	B
218	NA	NA	NA
219	B	B	B
220	NA	NA	NA
221	B	B	B
222	C	NA	NA
223	B	B	B
224	C	NA	NA
225	C	C	C

006090-19116560

226	A	A	B
227	NA	NA	NA
228	C	NA	NA
229	C	B	B
230	C	B	B
231	B	B	B
232	B	B	NA
233	B	B	C
234	B	B	B
235	B	B	B
236	B	B	B
237	B	A	B
238	B	B	B
239	C	B	B
240	C	B	B
241	B	B	B
242	B	B	B
243	B	B	B
244	B	B	A
245	B	B	B
246	B	B	B
247	B	B	B
248	B	B	B
249	B	B	B
250	B	B	B
251	C	B	B
252	B	B	B
253	C	B	B
254	B	B	B
255	C	NA	NA
256	C	NA	NA
257	NA	NA	NA

006090" inst 6560

258	C	NA	NA
259	C	B	B
260	B	B	B
261	B	B	B
262	B	B	B
263	B	B	B
264	C	C	NA
265	C	C	C
266	C	NA	NA
267	C	NA	NA
268	C	B	C
269	C	B	C
270	B	B	B
271	B	B	B
272	B	B	B
273	NA	NA	NA
274	C	B	C
275	C	B	C
276	NA	NA	NA
277	B	B	B
278	C	B	B
279	B	B	B
280	C	B	B
281	B	B	B
282	C	B	C
283	C	B	C
284	B	B	B
285	C	NA	NA
286	C	B	B
287	B	B	B
288	B	B	B
289	B	B	B

290	NA	NA	NA
291	NA	NA	NA
292	C	C	C
293	B	B	B
294	B	B	B
295	B	B	B
296	NA	NA	NA
297	C	C	NA
298	C	B	B
299	B	B	B
300	B	B	B
301	B	B	B
302	B	B	C
303	C	C	C
304	B	B	B
305	B	B	B
306	C	NA	NA
307	C	NA	NA
308	C	B	B
309	B	B	B
310	B	B	C
311	B	B	C
312	C	B	C
313	C	B	B
314	C	C	C
315	C	B	B
316	NA	NA	NA
317	B	B	B
318	B	B	B
319	B	B	B
320	B	B	B
321	B	B	B

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322	B	B	B
323	B	B	B
324	B	B	B
325	C	NA	NA
326	B	C	C
327	NA	NA	NA
328	C	NA	NA
329	B	C	C
330	C	NA	NA
331	C	C	C
332	B	B	C
333	B	B	B
334	B	B	B
335	B	B	B
336	B	B	B
337	B	B	B
338	B	B	B
339	C	NA	NA
340	C	NA	NA
341	NA	NA	NA
342	C	C	C
343	NA	NA	NA
344	NA	NA	NA
345	NA	NA	NA
346	C	NA	NA
347	B	B	B
348	B	B	B
349	B	B	B
350	C	NA	NA
351	NA	NA	NA
352	A	A	B
353	A	A	B

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354	C	NA	NA
355	C	C	C
356	B	B	A
357	C	B	C
358	C	C	C
359	C	C	C
360	NA	NA	NA
361	B	C	C
362	C	NA	NA
363	C	NA	NA
364	NA	NA	NA
365	B	B	B
366	B	B	B
367	A	A	B
368	A	A	B
369	B	NA	NA
370	A	B	B
371	C	B	C
372	B	B	C
373	B	B	C
374	B	B	B
375	B	B	C
376	B	C	C
377	B	B	C
378	B	B	C
379	B	B	B
380	B	B	C
381	B	B	B
382	B	B	B
383	B	B	B
384	B	B	B
385	B	B	C

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386	B	B	C
387	B	B	B
388	B	B	B
389	B	B	B
390	B	B	C
391	B	B	B
392	B	B	B
393	C	C	C
394	B	B	B
395	B	B	B
396	B	B	B
397	C	B	B
398	NA	NA	NA
399	NA	NA	NA
400	NA	NA	NA

In Table B, the following classifications have been employed:

A < 0.001 μ M

5 0.010 > B > 0.001 μ M

0.100 > C > 0.010 μ M

D > 0.1 μ M

"NA" = compound was not tested.

10 While we have hereinbefore presented a number of
embodiments of this invention, it is apparent that our
basic construction can be altered to provide other
embodiments which utilize the methods of this invention.
Therefore, it will be appreciated that the scope of this
15 invention is to be defined by the claims appended hereto
rather than the specific embodiments which have been
presented hereinbefore by way of example.